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(54) Title: CORYNEBACTERIUM GLUTAMICUM GENES ENCODING PROTEINS INVOLVED IN CARBON METABOLISM AND ENERGY PRODUCTION

(57) Abstract: Isolated nucleic acid molecules, designated SMP nucleic acid molecules, which encode novel SMP proteins from Corynebacterium glutamicum are described. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing SMP nucleic acid molecules, and host cells into which the expression vectors have been introduced. The invention still further provides isolated SMP proteins, mutated SMP proteins, fusion proteins, antigenic peptides and methods for the improvement of production of a desired compound from C. glutamicum based on genetic engineering of SMP genes in this organism.



CORYNEBACTERIUM GLUTAMICUM GENES ENCODING PROTEINS INVOLVED IN CARBON METABOLISM AND ENERGY PRODUCTION

Related Applications

5 This application claims priority to prior U.S. Provisional Patent Application Serial No. 60/141031, filed June 25, 1999, U.S. Provisional Patent Application Serial No. 60/143208, filed July 9, 1999, and U.S. Provisional Patent Application Serial No. 60/151572, filed August 31, 1999. This application also claims priority to prior German Patent Application No. 19931412.8, filed July 8, 1999, German Patent Application No. 19931413.6, filed July 8, 1999, German Patent Application No. 19931419.5, filed July 10 8, 1999, German Patent Application No. 19931420.9, filed July 8, 1999, German Patent Application No. 19931424.1, filed July 8, 1999, German Patent Application No. 19931428.4, filed July 8, 1999, German Patent Application No. 19931431.4, filed July 8, 1999, German Patent Application No. 19931433.0, filed July 8, 1999, German Patent Application No. 19931434.9, filed July 8, 1999, German Patent Application No. 15 19931510.8, filed July 8, 1999, German Patent Application No. 19931562.0, filed July 8, 1999, German Patent Application No. 19931634.1, filed July 8, 1999, German Patent Application No. 19932180.9, filed July 9, 1999, German Patent Application No. 19932227.9, filed July 9, 1999, German Patent Application No. 19932230.9, filed July 20 9, 1999, German Patent Application No. 19932924.9, filed July 14, 1999, German Patent Application No. 19932973.7, filed July 14, 1999, German Patent Application No. 19933005.0, filed July 14, 1999, German Patent Application No. 19940765.7, filed August 27, 1999, German Patent Application No. 19942076.9, filed September 3, 1999, German Patent Application No. 19942079.3, filed September 3, 1999, German Patent Application No. 19942086.6, filed September 3, 1999, German Patent Application No. 25 19942087.4, filed September 3, 1999, German Patent Application No. 19942088.2, filed September 3, 1999, German Patent Application No. 19942095.5, filed September 3, 1999, German Patent Application No. 19942123.4, filed September 3, 1999, and German Patent Application No. 19942125.0, filed September 3, 1999. The entire contents of all of the aforementioned application are hereby expressly incorporated herein by this reference.

Background of the Invention

Certain products and by-products of naturally-occurring metabolic processes in cells have utility in a wide array of industries, including the food, feed, cosmetics, and pharmaceutical industries. These molecules, collectively termed 'fine chemicals', include organic acids, both proteinogenic and non-proteinogenic amino acids, nucleotides and nucleosides, lipids and fatty acids, diols, carbohydrates, aromatic compounds, vitamins and cofactors, and enzymes. Their production is most conveniently performed through the large-scale culture of bacteria developed to produce and secrete large quantities of one or more desired molecules. One particularly useful organism for this purpose is *Corynebacterium glutamicum*, a gram positive, nonpathogenic bacterium. Through strain selection, a number of mutant strains have been developed which produce an array of desirable compounds. However, selection of strains improved for the production of a particular molecule is a time-consuming and difficult process.

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Summary of the Invention

The invention provides novel bacterial nucleic acid molecules which have a variety of uses. These uses include the identification of microorganisms which can be used to produce fine chemicals, the modulation of fine chemical production in C. glutamicum or related bacteria, the typing or identification of C. glutamicum or related bacteria, as reference points for mapping the C. glutamicum genome, and as markers for transformation. These novel nucleic acid molecules encode proteins, referred to herein as sugar metabolism and oxidative phosphorylation (SMP) proteins.

C. glutamicum is a gram positive, aerobic bacterium which is commonly used in industry for the large-scale production of a variety of fine chemicals, and also for the degradation of hydrocarbons (such as in petroleum spills) and for the oxidation of terpenoids. The SMP nucleic acid molecules of the invention, therefore, can be used to identify microorganisms which can be used to produce fine chemicals, e.g., by fermentation processes. Modulation of the expression of the SMP nucleic acids of the invention, or modification of the sequence of the SMP nucleic acid molecules of the invention, can be used to modulate the production of one or more fine chemicals from a

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microorganism (e.g., to improve the yield or production of one or more fine chemicals from a Corynebacterium or Brevibacterium species).

The SMP nucleic acids of the invention may also be used to identify an organism as being Corynebacterium glutamicum or a close relative thereof, or to identify the presence of C. glutamicum or a relative thereof in a mixed population of microorganisms. The invention provides the nucleic acid sequences of a number of C. glutamicum genes; by probing the extracted genomic DNA of a culture of a unique or mixed population of microorganisms under stringent conditions with a probe spanning a region of a C. glutamicum gene which is unique to this organism, one can ascertain whether this organism is present. Although Corynebacterium glutamicum itself is nonpathogenic, it is related to species pathogenic in humans, such as Corynebacterium diphtheriae (the causative agent of diphtheria); the detection of such organisms is of significant clinical relevance.

The SMP nucleic acid molecules of the invention may also serve as reference points for mapping of the *C. glutamicum* genome, or of genomes of related organisms. Similarly, these molecules, or variants or portions thereof, may serve as markers for genetically engineered Corynebacterium or Brevibacterium species.

The SMP proteins encoded by the novel nucleic acid molecules of the invention are capable of, for example, performing a function involved in the metabolism of carbon compounds such as sugars or in the generation of energy molecules by processes such as oxidative phosphorylation in *Corynebacterium glutamicum*. Given the availability of cloning vectors for use in *Corynebacterium glutamicum*, such as those disclosed in Sinskey *et al.*, U.S. Patent No. 4,649,119, and techniques for genetic manipulation of *C. glutamicum* and the related *Brevibacterium* species (*e.g.*, *lactofermentum*) (Yoshihama et al., *J. Bacteriol.* 162: 591-597 (1985); Katsumata *et al.*, *J. Bacteriol.* 159: 306-311 (1984); and Santamaria *et al.*, *J. Gen. Microbiol.* 130: 2237-2246 (1984)), the nucleic acid molecules of the invention may be utilized in the genetic engineering of this organism to make it a better or more efficient producer of one or more fine chemicals. This improved production or efficiency of production of a fine chemical may be due to a direct effect of manipulation of a gene of the invention, or it may be due to an indirect effect of such manipulation.

WO 01/00844 PCT/IB00/00943

- 4 -

There are a number of mechanisms by which the alteration of an SMP protein of the invention may directly affect the yield, production, and/or efficiency of production of a fine chemical from a C. glutamicum strain incorporating such an altered protein. The degradation of high-energy carbon molecules such as sugars, and the conversion of compounds such as NADH and FADH2 to compounds containing high energy phosphate bonds via oxidative phosphorylation results in a number of compounds which themselves may be desirable fine chemicals, such as pyruvate, ATP, NADH, and a number of intermediate sugar compounds. Further, the energy molecules (such as ATP) and the reducing equivalents (such as NADH or NADPH) produced by these metabolic pathways are utilized in the cell to drive reactions which would otherwise be energetically unfavorable. Such unfavorable reactions include many biosynthetic pathways for fine chemicals. By improving the ability of the cell to utilize a particular sugar (e.g., by manipulating the genes encoding enzymes involved in the degradation and conversion of that sugar into energy for the cell), one may increase the amount of energy available to permit unfavorable, yet desired metabolic reactions (e.g., the biosynthesis of a desired fine chemical) to occur.

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The mutagenesis of one or more SMP genes of the invention may also result in SMP proteins having altered activities which indirectly impact the production of one or more desired fine chemicals from C. glutamicum. For example, by increasing the efficiency of utilization of one or more sugars (such that the conversion of the sugar to useful energy molecules is improved), or by increasing the efficiency of conversion of reducing equivalents to useful energy molecules (e.g., by improving the efficiency of oxidative phosphorylation, or the activity of the ATP synthase), one can increase the amount of these high-energy compounds available to the cell to drive normally unfavorable metabolic processes. These processes include the construction of cell walls, transcription, translation, and the biosynthesis of compounds necessary for growth and division of the cells (e.g., nucleotides, amino acids, vitamins, lipids, etc.) (Lengeler et al. (1999) Biology of Prokaryotes, Thieme Verlag: Stuttgart, p. 88-109; 913-918; 875-899). By improving the growth and multiplication of these engineered cells, it is possible to increase both the viability of the cells in large-scale culture, and also to improve their rate of division, such that a relatively larger number of cells can survive in fermentor culture. The yield, production, or efficiency of production may be increased, at least

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due to the presence of a greater number of viable cells, each producing the desired fine chemical. Also, many of the degradation products produced during sugar metabolism are utilized by the cell as precursors or intermediates in the production of other desirable products, such as fine chemicals. So, by increasing the ability of the cell to metabolize sugars, the number of these degradation products available to the cell for other processes should also be increased.

The invention provides novel nucleic acid molecules which encode proteins, referred to herein as SMP proteins, which are capable of, for example, performing a function involved in the metabolism of carbon compounds such as sugars and the generation of energy molecules by processes such as oxidative phosphorylation in *Corynebacterium glutamicum*. Nucleic acid molecules encoding an SMP protein are referred to herein as SMP nucleic acid molecules. In a preferred embodiment, the SMP protein participates in the conversion of carbon molecules and degradation products thereof to energy which is utilized by the cell for metabolic processes. Examples of such proteins include those encoded by the genes set forth in Table 1.

Accordingly, one aspect of the invention pertains to isolated nucleic acid molecules (e.g., cDNAs, DNAs, or RNAs) comprising a nucleotide sequence encoding an SMP protein or biologically active portions thereof, as well as nucleic acid fragments suitable as primers or hybridization probes for the detection or amplification of SMPencoding nucleic acid (e.g., DNA or mRNA). In particularly preferred embodiments, the isolated nucleic acid molecule comprises one of the nucleotide sequences set forth as the odd-numbered SEQ ID NOs in the Sequence Listing (e.g., SEQ ID NO:1, SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7....), or the coding region or a complement thereof of one of these nucleotide sequences. In other particularly preferred embodiments, the isolated nucleic acid molecule of the invention comprises a nucleotide sequence which hybridizes to or is at least about 50%, preferably at least about 60%, more preferably at least about 70%, 80% or 90%, and even more preferably at least about 95%, 96%, 97%, 98%, 99% or more homologous to a nucleotide sequence set forth as an odd-numbered SEQ ID NO in the Sequence Listing (e.g., SEQ ID NO:1, SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7....), or a portion thereof. In other preferred embodiments, the isolated nucleic acid molecule encodes one of the amino acid sequences set forth as an evennumbered SEQ ID NO in the Sequence Listing (e.g., SEQ ID NO:2, SEQ ID NO:4, SEQ

ID NO:6, SEQ ID NO:8....).. The preferred SMP proteins of the present invention also preferably possess at least one of the SMP activities described herein.

In another embodiment, the isolated nucleic acid molecule encodes a protein or portion thereof wherein the protein or portion thereof includes an amino acid sequence which is sufficiently homologous to an amino acid sequence of the invention (e.g., a sequence having an even-numbered SEQ ID NO: in the Sequence Listing), e.g., sufficiently homologous to an amino acid sequence of the invention such that the protein or portion thereof maintains an SMP activity. Preferably, the protein or portion thereof encoded by the nucleic acid molecule maintains the ability to perform a function involved in the metabolism of carbon compounds such as sugars or the generation of energy molecules (e.g., ATP) by processes such as oxidative phosphorylation in Corynebacterium glutamicum. In one embodiment, the protein encoded by the nucleic acid molecule is at least about 50%, preferably at least about 60%, and more preferably at least about 70%, 80%, or 90% and most preferably at least about 95%, 96%, 97%, 98%, or 99% or more homologous to an amino acid sequence of the invention (e.g., an entire amino acid sequence selected those having an even-numbered SEQ ID NO in the Sequence Listing). In another preferred embodiment, the protein is a full length C. glutamicum protein which is substantially homologous to an entire amino acid sequence of the invention (encoded by an open reading frame shown in the corresponding oddnumbered SEQ ID NOs in the Sequence Listing (e.g., SEQ ID NO:1, SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7....).

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In another preferred embodiment, the isolated nucleic acid molecule is derived from *C. glutamicum* and encodes a protein (*e.g.*, an SMP fusion protein) which includes a biologically active domain which is at least about 50% or more homologous to one of the amino acid sequences of the invention (*e.g.*, a sequence of one of the even-numbered SEQ ID NOs in the Sequence Listing) and is able to perform a function involved in the metabolism of carbon compounds such as sugars or the generation of energy molecules (*e.g.*, ATP) by processes such as oxidative phosphorylation in *Corynebacterium glutamicum*, or has one or more of the activities set forth in Table 1, and which also includes heterologous nucleic acid sequences encoding a heterologous polypeptide or regulatory regions.

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In another embodiment, the isolated nucleic acid molecule is at least 15 nucleotides in length and hybridizes under stringent conditions to a nucleic acid molecule comprising a nucleotide sequence of the invention (e.g., a sequence of an odd-numbered SEQ ID NO in the Sequence Listing) A. Preferably, the isolated nucleic acid molecule corresponds to a naturally-occurring nucleic acid molecule. More preferably, the isolated nucleic acid encodes a naturally-occurring C. glutamicum SMP protein, or a biologically active portion thereof.

Another aspect of the invention pertains to vectors, e.g., recombinant expression vectors, containing the nucleic acid molecules of the invention, and host cells into which such vectors have been introduced. In one embodiment, such a host cell is used to produce an SMP protein by culturing the host cell in a suitable medium. The SMP protein can be then isolated from the medium or the host cell.

Yet another aspect of the invention pertains to a genetically altered microorganism in which an SMP gene has been introduced or altered. In one embodiment, the genome of the microorganism has been altered by introduction of a nucleic acid molecule of the invention encoding wild-type or mutated SMP sequence as a transgene. In another embodiment, an endogenous SMP gene within the genome of the microorganism has been altered, e.g., functionally disrupted, by homologous recombination with an altered SMP gene. In another embodiment, an endogenous or introduced SMP gene in a microorganism has been altered by one or more point mutations, deletions, or inversions, but still encodes a functional SMP protein. In still another embodiment, one or more of the regulatory regions (e.g., a promoter, repressor, or inducer) of an SMP gene in a microorganism has been altered (e.g., by deletion, truncation, inversion, or point mutation) such that the expression of the SMP gene is modulated. In a preferred embodiment, the microorganism belongs to the genus Corynebacterium or Brevibacterium, with Corynebacterium glutamicum being particularly preferred. In a preferred embodiment, the microorganism is also utilized for the production of a desired compound, such as an amino acid, with lysine being particularly preferred.

In another aspect, the invention provides a method of identifying the presence or activity of *Cornyebacterium diphtheriae* in a subject. This method includes detection of one or more of the nucleic acid or amino acid sequences of the invention (e.g., the

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sequences set forth in the Sequence Listing as SEQ ID NOs 1 through 782) in a subject, thereby detecting the presence or activity of *Corynebacterium diphtheriae* in the subject.

Still another aspect of the invention pertains to an isolated SMP protein or a portion, e.g., a biologically active portion, thereof. In a preferred embodiment, the isolated SMP protein or portion thereof is capable of performing a function involved in the metabolism of carbon compounds such as sugars or in the generation of energy molecules (e.g., ATP) by processes such as oxidative phosphorylation in Corynebacterium glutamicum. In another preferred embodiment, the isolated SMP protein or portion thereof is sufficiently homologous to an amino acid sequence of the invention (e.g., a sequence of an even-numbered SEQ ID NO: in the Sequence Listing) such that the protein or portion thereof maintains the ability to perform a function involved in the metabolism of carbon compounds such as sugars or in the generation of energy molecules (e.g., ATP) by processes such as oxidative phosphorylation in Corynebacterium glutamicum.

The invention also provides an isolated preparation of an SMP protein. In preferred embodiments, the SMP protein comprises an amino acid sequence of the invention (e.g., a sequence of an even-numbered SEQ ID NO: of the Sequence Listing). In another preferred embodiment, the invention pertains to an isolated full length protein which is substantially homologous to an entire amino acid sequence of the invention (e.g., a sequence of an even-numbered SEQ ID NO: of the Sequence Listing) (encoded by an open reading frame set forth in a corresponding odd-numbered SEQ ID NO: of the Sequence Listing). In yet another embodiment, the protein is at least about 50%, preferably at least about 60%, and more preferably at least about 70%, 80%, or 90%, and most preferably at least about 95%, 96%, 97%, 98%, or 99% or more homologous to an entire amino acid sequence of the invention (e.g., a sequence of an even-numbered SEQ ID NO: of the Sequence Listing). In other embodiments, the isolated SMP protein comprises an amino acid sequence which is at least about 50% or more homologous to one of the amino acid sequences of the invention (e.g., a sequence of an even-numbered SEQ ID NO: of the Sequence Listing) and is able to perform a function involved in the metabolism of carbon compounds such as sugars or in the generation of energy molecules (e.g., ATP) by processes such as oxidative phosphorylation in Corynebacterium glutamicum, or has one or more of the activities set forth in Table 1.

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Alternatively, the isolated SMP protein can comprise an amino acid sequence which is encoded by a nucleotide sequence which hybridizes, e.g., hybridizes under stringent conditions, or is at least about 50%, preferably at least about 60%, more preferably at least about 70%, 80%, or 90%, and even more preferably at least about 95%, 96%, 97%, 98,%, or 99% or more homologous to a nucleotide sequence of one of the even-numbered SEQ ID NOs set forth in the Sequence Listing. It is also preferred that the preferred forms of SMP proteins also have one or more of the SMP bioactivities described herein.

10 linked to a non-SMP polypeptide to form a fusion protein. In preferred embodiments, this fusion protein has an activity which differs from that of the SMP protein alone. In other preferred embodiments, this fusion protein performs a function involved in the metabolism of carbon compounds such as sugars or in the generation of energy molecules (e.g., ATP) by processes such as oxidative phosphorylation in

15 Corynebacterium glutamicum. In particularly preferred embodiments, integration of this fusion protein into a host cell modulates production of a desired compound from the cell.

In another aspect, the invention provides methods for screening molecules which modulate the activity of an SMP protein, either by interacting with the protein itself or a substrate or binding partner of the SMP protein, or by modulating the transcription or translation of an SMP nucleic acid molecule of the invention.

Another aspect of the invention pertains to a method for producing a fine chemical. This method involves the culturing of a cell containing a vector directing the expression of an SMP nucleic acid molecule of the invention, such that a fine chemical is produced. In a preferred embodiment, this method further includes the step of obtaining a cell containing such a vector, in which a cell is transfected with a vector directing the expression of an SMP nucleic acid. In another preferred embodiment, this method further includes the step of recovering the fine chemical from the culture. In a particularly preferred embodiment, the cell is from the genus *Corynebacterium* or *Brevibacterium*, or is selected from those strains set forth in Table 3.

Another aspect of the invention pertains to methods for modulating production of a molecule from a microorganism. Such methods include contacting the cell with an

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agent which modulates SMP protein activity or SMP nucleic acid expression such that a cell associated activity is altered relative to this same activity in the absence of the agent. In a preferred embodiment, the cell is modulated for one or more *C. glutamicum* carbon metabolism pathways or for the production of energy through processes such as oxidative phosphorylation, such that the yields or rate of production of a desired fine chemical by this microorganism is improved. The agent which modulates SMP protein activity can be an agent which stimulates SMP protein activity or SMP nucleic acid expression. Examples of agents which stimulate SMP proteins, and nucleic acids encoding SMP proteins that have been introduced into the cell. Examples of agents which inhibit SMP activity or expression include small molecules and antisense SMP nucleic acid molecules.

Another aspect of the invention pertains to methods for modulating yields of a desired compound from a cell, involving the introduction of a wild-type or mutant SMP gene into a cell, either maintained on a separate plasmid or integrated into the genome of the host cell. If integrated into the genome, such integration can be random, or it can take place by homologous recombination such that the native gene is replaced by the introduced copy, causing the production of the desired compound from the cell to be modulated. In a preferred embodiment, said yields are increased. In another preferred embodiment, said chemical is a fine chemical. In a particularly preferred embodiment, said fine chemical is an amino acid. In especially preferred embodiments, said amino acid is L-lysine.

Detailed Description of the Invention

The present invention provides SMP nucleic acid and protein molecules which are involved in the metabolism of carbon compounds such as sugars and the generation of energy molecules by processes such as oxidative phosphorylation in Corynebacterium glutamicum. The molecules of the invention may be utilized in the modulation of production of fine chemicals from microorganisms, such as C. glutamicum, either directly (e.g., where overexpression or optimization of a glycolytic pathway protein has a direct impact on the yield, production, and/or efficiency of production of, e.g., pyruvate from modified C. glutamicum), or may have an indirect

impact which nonetheless results in an increase of yield, production, and/or efficiency of production of the desired compound (e.g., where modulation of proteins involved in oxidative phosphorylation results in alterations in the amount of energy available to perform necessary metabolic processes and other cellular functions, such as nucleic acid and protein biosynthesis and transcription/translation). Aspects of the invention are further explicated below.

I. Fine Chemicals

The term 'fine chemical' is art-recognized and includes molecules produced by 10 an organism which have applications in various industries, such as, but not limited to, the pharmaceutical, agriculture, and cosmetics industries. Such compounds include organic acids, such as tartaric acid, itaconic acid, and diaminopimelic acid, both proteinogenic and non-proteinogenic amino acids, purine and pyrimidine bases, nucleosides, and nucleotides (as described e.g. in Kuninaka, A. (1996) Nucleotides and 15 related compounds, p. 561-612, in Biotechnology vol. 6, Rehm et al., eds. VCH: Weinheim, and references contained therein), lipids, both saturated and unsaturated fatty acids (e.g., arachidonic acid), diols (e.g., propane diol, and butane diol), carbohydrates (e.g., hyaluronic acid and trehalose), aromatic compounds (e.g., aromatic amines, vanillin, and indigo), vitamins and cofactors (as described in Ullmann's Encyclopedia of Industrial Chemistry, vol. A27, "Vitamins", p. 443-613 (1996) VCH: Weinheim and 20 references therein; and Ong, A.S., Niki, E. & Packer, L. (1995) "Nutrition, Lipids, Health, and Disease" Proceedings of the UNESCO/Confederation of Scientific and Technological Associations in Malaysia, and the Society for Free Radical Research -Asia, held Sept. 1-3, 1994 at Penang, Malaysia, AOCS Press, (1995)), enzymes, polyketides (Cane et al. (1998) Science 282: 63-68), and all other chemicals described in 25 Gutcho (1983) Chemicals by Fermentation, Noyes Data Corporation, ISBN: 0818805086 and references therein. The metabolism and uses of certain of these fine chemicals are further explicated below.

30 A. Amino Acid Metabolism and Uses

Amino acids comprise the basic structural units of all proteins, and as such are essential for normal cellular functioning in all organisms. The term "amino acid" is art-

WO 01/00844 PCT/IB00/00943

- 12 -

recognized. The proteinogenic amino acids, of which there are 20 species, serve as structural units for proteins, in which they are linked by peptide bonds, while the nonproteinogenic amino acids (hundreds of which are known) are not normally found in proteins (see Ulmann's Encyclopedia of Industrial Chemistry, vol. A2, p. 57-97 VCH: Weinheim (1985)). Amino acids may be in the D- or L- optical configuration, though Lamino acids are generally the only type found in naturally-occurring proteins. Biosynthetic and degradative pathways of each of the 20 proteinogenic amino acids have been well characterized in both prokaryotic and eukaryotic cells (see, for example, Stryer, L. Biochemistry, 3rd edition, pages 578-590 (1988)). The 'essential' amino acids (histidine, isoleucine, leucine, lysine, methionine, phenylalanine, threonine, tryptophan, 10 and valine), so named because they are generally a nutritional requirement due to the complexity of their biosyntheses, are readily converted by simple biosynthetic pathways to the remaining 11 'nonessential' amino acids (alanine, arginine, asparagine, aspartate, cysteine, glutamate, glutamine, glycine, proline, serine, and tyrosine). Higher animals do retain the ability to synthesize some of these amino acids, but the essential amino 15 acids must be supplied from the diet in order for normal protein synthesis to occur.

Aside from their function in protein biosynthesis, these amino acids are interesting chemicals in their own right, and many have been found to have various applications in the food, feed, chemical, cosmetics, agriculture, and pharmaceutical industries. Lysine is an important amino acid in the nutrition not only of humans, but also of monogastric animals such as poultry and swine. Glutamate is most commonly used as a flavor additive (mono-sodium glutamate, MSG) and is widely used throughout the food industry, as are aspartate, phenylalanine, glycine, and cysteine. Glycine, L-methionine and tryptophan are all utilized in the pharmaceutical industry. Glutamine, valine, leucine, isoleucine, histidine, arginine, proline, serine and alanine are of use in both the pharmaceutical and cosmetics industries. Threonine, tryptophan, and D/L-methionine are common feed additives. (Leuchtenberger, W. (1996) Amino aids – technical production and use, p. 466-502 in Rehm *et al.* (eds.) Biotechnology vol. 6, chapter 14a, VCH: Weinheim). Additionally, these amino acids have been found to be useful as precursors for the synthesis of synthetic amino acids and proteins, such as N-acetylcysteine, S-carboxymethyl-L-cysteine, (S)-5-hydroxytryptophan, and others

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described in Ulmann's Encyclopedia of Industrial Chemistry, vol. A2, p. 57-97, VCH: Weinheim, 1985.

The biosynthesis of these natural amino acids in organisms capable of producing them, such as bacteria, has been well characterized (for review of bacterial amino acid biosynthesis and regulation thereof, see Umbarger, H.E.(1978) Ann. Rev. Biochem. 47: 533-606). Glutamate is synthesized by the reductive amination of αketoglutarate, an intermediate in the citric acid cycle. Glutamine, proline, and arginine are each subsequently produced from glutamate. The biosynthesis of serine is a threestep process beginning with 3-phosphoglycerate (an intermediate in glycolysis), and resulting in this amino acid after oxidation, transamination, and hydrolysis steps. Both cysteine and glycine are produced from serine; the former by the condensation of homocysteine with serine, and the latter by the transferal of the side-chain β-carbon atom to tetrahydrofolate, in a reaction catalyzed by serine transhydroxymethylase. Phenylalanine, and tyrosine are synthesized from the glycolytic and pentose phosphate pathway precursors erythrose 4-phosphate and phosphoenolpyruvate in a 9-step biosynthetic pathway that differ only at the final two steps after synthesis of prephenate. Tryptophan is also produced from these two initial molecules, but its synthesis is an 11step pathway. Tyrosine may also be synthesized from phenylalanine, in a reaction catalyzed by phenylalanine hydroxylase. Alanine, valine, and leucine are all biosynthetic products of pyruvate, the final product of glycolysis. Aspartate is formed from oxaloacetate, an intermediate of the citric acid cycle. Asparagine, methionine, threonine, and lysine are each produced by the conversion of aspartate. Isoleucine is formed from threonine. A complex 9-step pathway results in the production of histidine from 5-phosphoribosyl-1-pyrophosphate, an activated sugar.

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Amino acids in excess of the protein synthesis needs of the cell cannot be stored, and are instead degraded to provide intermediates for the major metabolic pathways of the cell (for review see Stryer, L. Biochemistry 3rd ed. Ch. 21 "Amino Acid Degradation and the Urea Cycle" p. 495-516 (1988)). Although the cell is able to convert unwanted amino acids into useful metabolic intermediates, amino acid production is costly in terms of energy, precursor molecules, and the enzymes necessary to synthesize them. Thus it is not surprising that amino acid biosynthesis is regulated by feedback inhibition, in which the presence of a particular amino acid serves to slow or entirely stop its own

production (for overview of feedback mechanisms in amino acid biosynthetic pathways, see Stryer, L. Biochemistry, 3rd ed. Ch. 24: "Biosynthesis of Amino Acids and Heme" p. 575-600 (1988)). Thus, the output of any particular amino acid is limited by the amount of that amino acid present in the cell.

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B. Vitamin, Cofactor, and Nutraceutical Metabolism and Uses

Vitamins, cofactors, and nutraceuticals comprise another group of molecules which the higher animals have lost the ability to synthesize and so must ingest, although they are readily synthesized by other organisms such as bacteria. These molecules are either bioactive substances themselves, or are precursors of biologically active substances which may serve as electron carriers or intermediates in a variety of metabolic pathways. Aside from their nutritive value, these compounds also have significant industrial value as coloring agents, antioxidants, and catalysts or other processing aids. (For an overview of the structure, activity, and industrial applications of these compounds, see, for example, Ullman's Encyclopedia of Industrial Chemistry, "Vitamins" vol. A27, p. 443-613, VCH: Weinheim, 1996.) The term "vitamin" is artrecognized, and includes nutrients which are required by an organism for normal functioning, but which that organism cannot synthesize by itself. The group of vitamins may encompass cofactors and nutraceutical compounds. The language "cofactor" includes nonproteinaceous compounds required for a normal enzymatic activity to occur. Such compounds may be organic or inorganic; the cofactor molecules of the invention are preferably organic. The term "nutraceutical" includes dietary supplements having health benefits in plants and animals, particularly humans. Examples of such molecules are vitamins, antioxidants, and also certain lipids (e.g., polyunsaturated fatty acids).

The biosynthesis of these molecules in organisms capable of producing them, such as bacteria, has been largely characterized (Ullman's Encyclopedia of Industrial Chemistry, "Vitamins" vol. A27, p. 443-613, VCH: Weinheim, 1996; Michal, G. (1999) Biochemical Pathways: An Atlas of Biochemistry and Molecular Biology, John Wiley & Sons; Ong, A.S., Niki, E. & Packer, L. (1995) "Nutrition, Lipids, Health, and Disease" Proceedings of the UNESCO/Confederation of Scientific and Technological

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Associations in Malaysia, and the Society for Free Radical Research - Asia, held Sept. 1-3, 1994 at Penang, Malaysia, AOCS Press: Champaign, IL X, 374 S).

Thiamin (vitamin B₁) is produced by the chemical coupling of pyrimidine and thiazole moieties. Riboflavin (vitamin B2) is synthesized from guanosine-5'-triphosphate (GTP) and ribose-5'-phosphate. Riboflavin, in turn, is utilized for the synthesis of flavin mononucleotide (FMN) and flavin adenine dinucleotide (FAD). The family of compounds collectively termed 'vitamin B₆' (e.g., pyridoxine, pyridoxamine, pyridoxa-5'-phosphate, and the commercially used pyridoxin hydrochloride) are all derivatives of the common structural unit, 5-hydroxy-6-methylpyridine. Pantothenate (pantothenic acid, (R)-(+)-N-(2,4-dihydroxy-3,3-dimethyl-1-oxobutyl)-β-alanine) can be produced either by chemical synthesis or by fermentation. The final steps in pantothenate biosynthesis consist of the ATP-driven condensation of \beta-alanine and pantoic acid. The enzymes responsible for the biosynthesis steps for the conversion to pantoic acid, to \betaalanine and for the condensation to panthotenic acid are known. The metabolically active form of pantothenate is Coenzyme A, for which the biosynthesis proceeds in 5 enzymatic steps. Pantothenate, pyridoxal-5'-phosphate, cysteine and ATP are the precursors of Coenzyme A. These enzymes not only catalyze the formation of panthothante, but also the production of (R)-pantoic acid, (R)-pantolacton, (R)panthenol (provitamin B₅), pantetheine (and its derivatives) and coenzyme A.

20 Biotin biosynthesis from the precursor molecule pimeloyl-CoA in microorganisms has been studied in detail and several of the genes involved have been identified. Many of the corresponding proteins have been found to also be involved in Fe-cluster synthesis and are members of the nifS class of proteins. Lipoic acid is derived from octanoic acid, and serves as a coenzyme in energy metabolism, where it becomes part of the pyruvate dehydrogenase complex and the α-ketoglutarate dehydrogenase complex. The folates are a group of substances which are all derivatives of folic acid, which is turn is derived from L-glutamic acid, p-amino-benzoic acid and 6methylpterin. The biosynthesis of folic acid and its derivatives, starting from the metabolism intermediates guanosine-5'-triphosphate (GTP), L-glutamic acid and pamino-benzoic acid has been studied in detail in certain microorganisms.

Corrinoids (such as the cobalamines and particularly vitamin B₁₂) and porphyrines belong to a group of chemicals characterized by a tetrapyrole ring system.

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The biosynthesis of vitamin B₁₂ is sufficiently complex that it has not yet been completely characterized, but many of the enzymes and substrates involved are now known. Nicotinic acid (nicotinate), and nicotinamide are pyridine derivatives which are also termed 'niacin'. Niacin is the precursor of the important coenzymes NAD (nicotinamide adenine dinucleotide) and NADP (nicotinamide adenine dinucleotide phosphate) and their reduced forms.

The large-scale production of these compounds has largely relied on cell-free chemical syntheses, though some of these chemicals have also been produced by large-scale culture of microorganisms, such as riboflavin, Vitamin B₆, pantothenate, and biotin. Only Vitamin B₁₂ is produced solely by fermentation, due to the complexity of its synthesis. *In vitro* methodologies require significant inputs of materials and time, often at great cost.

C. Purine, Pyrimidine, Nucleoside and Nucleotide Metabolism and Uses

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Purine and pyrimidine metabolism genes and their corresponding proteins are important targets for the therapy of tumor diseases and viral infections. The language "purine" or "pyrimidine" includes the nitrogenous bases which are constituents of nucleic acids, co-enzymes, and nucleotides. The term "nucleotide" includes the basic structural units of nucleic acid molecules, which are comprised of a nitrogenous base, a pentose sugar (in the case of RNA, the sugar is ribose; in the case of DNA, the sugar is D-deoxyribose), and phosphoric acid. The language "nucleoside" includes molecules which serve as precursors to nucleotides, but which are lacking the phosphoric acid moiety that nucleotides possess. By inhibiting the biosynthesis of these molecules, or their mobilization to form nucleic acid molecules, it is possible to inhibit RNA and DNA synthesis; by inhibiting this activity in a fashion targeted to cancerous cells, the ability of tumor cells to divide and replicate may be inhibited. Additionally, there are nucleotides which do not form nucleic acid molecules, but rather serve as energy stores (i.e., AMP) or as coenzymes (i.e., FAD and NAD).

Several publications have described the use of these chemicals for these medical indications, by influencing purine and/or pyrimidine metabolism (e.g. Christopherson, R.I. and Lyons, S.D. (1990) "Potent inhibitors of de novo pyrimidine and purine biosynthesis as chemotherapeutic agents." Med. Res. Reviews 10: 505-548). Studies of

enzymes involved in purine and pyrimidine metabolism have been focused on the development of new drugs which can be used, for example, as immunosuppressants or anti-proliferants (Smith, J.L., (1995) "Enzymes in nucleotide synthesis." *Curr. Opin. Struct. Biol.* 5: 752-757; (1995) *Biochem Soc. Transact.* 23: 877-902). However, purine and pyrimidine bases, nucleosides and nucleotides have other utilities: as intermediates in the biosynthesis of several fine chemicals (*e.g.*, thiamine, S-adenosyl-methionine, folates, or riboflavin), as energy carriers for the cell (*e.g.*, ATP or GTP), and for chemicals themselves, commonly used as flavor enhancers (*e.g.*, IMP or GMP) or for several medicinal applications (see, for example, Kuninaka, A. (1996) Nucleotides and Related Compounds in Biotechnology vol. 6, Rehm *et al.*, eds. VCH: Weinheim, p. 561-612). Also, enzymes involved in purine, pyrimidine, nucleoside, or nucleotide metabolism are increasingly serving as targets against which chemicals for crop protection, including fungicides, herbicides and insecticides, are developed.

The metabolism of these compounds in bacteria has been characterized (for 15 reviews see, for example, Zalkin, H. and Dixon, J.E. (1992) "de novo purine nucleotide biosynthesis", in: Progress in Nucleic Acid Research and Molecular Biology, vol. 42, Academic Press:, p. 259-287; and Michal, G. (1999) "Nucleotides and Nucleosides", Chapter 8 in: Biochemical Pathways: An Atlas of Biochemistry and Molecular Biology, Wiley: New York). Purine metabolism has been the subject of intensive research, and is essential to the normal functioning of the cell. Impaired purine metabolism in higher 20 animals can cause severe disease, such as gout. Purine nucleotides are synthesized from ribose-5-phosphate, in a series of steps through the intermediate compound inosine-5'phosphate (IMP), resulting in the production of guanosine-5'-monophosphate (GMP) or adenosine-5'-monophosphate (AMP), from which the triphosphate forms utilized as nucleotides are readily formed. These compounds are also utilized as energy stores, so 25 their degradation provides energy for many different biochemical processes in the cell. Pyrimidine biosynthesis proceeds by the formation of uridine-5'-monophosphate (UMP) from ribose-5-phosphate. UMP, in turn, is converted to cytidine-5'-triphosphate (CTP). The deoxy- forms of all of these nucleotides are produced in a one step reduction reaction from the diphosphate ribose form of the nucleotide to the diphosphate 30 deoxyribose form of the nucleotide. Upon phosphorylation, these molecules are able to participate in DNA synthesis.

D. Trehalose Metabolism and Uses

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Trehalose consists of two glucose molecules, bound in α, α-1,1 linkage. It is commonly used in the food industry as a sweetener, an additive for dried or frozen foods, and in beverages. However, it also has applications in the pharmaceutical, cosmetics and biotechnology industries (see, for example, Nishimoto *et al.*, (1998) U.S. Patent No. 5,759,610; Singer, M.A. and Lindquist, S. (1998) *Trends Biotech.* 16: 460-467; Paiva, C.L.A. and Panek, A.D. (1996) *Biotech. Ann. Rev.* 2: 293-314; and Shiosaka, M. (1997) J. Japan 172: 97-102). Trehalose is produced by enzymes from many microorganisms and is naturally released into the surrounding medium, from which it can be collected using methods known in the art.

II. Sugar and Carbon Molecule Utilization and Oxidative Phosphorylation

Carbon is a critically important element for the formation of all organic compounds, and thus is a nutritional requirement not only for the growth and division of C. glutamicum, but also for the overproduction of fine chemicals from this microorganism. Sugars, such as mono-, di-, or polysaccharides, are particularly good carbon sources, and thus standard growth media typically contain one or more of: glucose, fructose, mannose, galactose, ribose, sorbose, ribulose, lactose, maltose, sucrose, raffinose, starch, or cellulose (Ullmann's Encyclopedia of Industrial Chemistry (1987) vol. A9, "Enzymes", VCH: Weinheim). Alternatively, more complex forms of sugar may be utilized in the media, such as molasses, or other by-products of sugar refinement. Other compounds aside from the sugars may be used as alternate carbon sources, including alcohols (e.g., ethanol or methanol), alkanes, sugar alcohols, fatty acids, and organic acids (e.g., acetic acid or lactic acid). For a review of carbon sources and their utilization by microorganisms in culture, see: Ullman's Encyclopedia of Industrial Chemistry (1987) vol. A9, "Enzymes", VCH: Weinheim; Stoppok, E. and Buchholz, K. (1996) "Sugar-based raw materials for fermentation applications" in Biotechnology (Rehm, H.J. et al., eds.) vol. 6, VCH: Weinheim, p. 5-29; Rehm, H.J. (1980) Industrielle Mikrobiologie, Springer: Berlin; Bartholomew, W.H., and Reiman, H.B. (1979). Economics of Fermentation Processes, in: Peppler, H.J. and Perlman, D., eds. Microbial Technology 2nd ed., vol. 2, chapter 18, Academic Press: New York; and

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Kockova-Kratachvilova, A. (1981) Characteristics of Industrial Microorganisms, in: Rehm, H.J. and Reed, G., eds. Handbook of Biotechnology, vol. 1, chapter 1, Verlag Chemie: Weinheim.

After uptake, these energy-rich carbon molecules must be processed such that

they are able to be degraded by one of the major sugar metabolic pathways. Such
pathways lead directly to useful degradation products, such as ribose-5-phosphate and
phosphoenolpyruvate, which may be subsequently converted to pyruvate. Three of the
most important pathways in bacteria for sugar metabolism include the EmbdenMeyerhoff-Pamas (EMP) pathway (also known as the glycolytic or fructose

bisphosphate pathway), the hexosemonophosphate (HMP) pathway (also known as the
pentose shunt or pentose phosphate pathway), and the Entner-Doudoroff (ED) pathway
(for review, see Michal, G. (1999) Biochemical Pathways: An Atlas of Biochemistry
and Molecular Biology, Wiley: New York, and Stryer, L. (1988) Biochemistry, Chapters
13-19, Freeman: New York, and references therein).

The EMP pathway converts hexose molecules to pyruvate, and in the process produces 2 molecules of ATP and 2 molecules of NADH. Starting with glucose-1-phosphate (which may be either directly taken up from the medium, or alternatively may be generated from glycogen, starch, or cellulose), the glucose molecule is isomerized to fructose-6-phosphate, is phosphorylated, and split into two 3-carbon molecules of glyceraldehyde-3-phosphate. After dehydrogenation, phosphorylation, and successive rearrangements, pyruvate results.

The HMP pathway converts glucose to reducing equivalents, such as NADPH, and produces pentose and tetrose compounds which are necessary as intermediates and precursors in a number of other metabolic pathways. In the HMP pathway, glucose-6-phosphate is converted to ribulose-5-phosphate by two successive dehydrogenase reactions (which also release two NADPH molecules), and a carboxylation step. Ribulose-5-phosphate may also be converted to xyulose-5-phosphate and ribose-5-phosphate; the former can undergo a series of biochemical steps to glucose-6-phosphate, which may enter the EMP pathway, while the latter is commonly utilized as an intermediate in other biosynthetic pathways within the cell.

The ED pathway begins with the compound glucose or gluconate, which is subsequently phosphorylated and dehydrated to form 2-dehydro-3-deoxy-6-P-gluconate.

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Glucuronate and galacturonate may also be converted to 2-dehydro-3-deoxy-6-P-gluconate through more complex biochemical pathways. This product molecule is subsequently cleaved into glyceraldehyde-3-P and pyruvate; glyceraldehyde-3-P may itself also be converted to pyruvate.

The EMP and HMP pathways share many features, including intermediates and enzymes. The EMP pathway provides the greatest amount of ATP, but it does not produce ribose-5-phosphate, an important precursor for, e.g., nucleic acid biosynthesis, nor does it produce erythrose-4-phosphate, which is important for amino acid biosynthesis. Microorganisms that are capable of using only the EMP pathway for glucose utilization are thus not able to grow on simple media with glucose as the sole carbon source. They are referred to as fastidious organisms, and their growth requires inputs of complex organic compounds, such as those found in yeast extract.

In contrast, the HMP pathway produces all of the precursors necessary for both nucleic acid and amino acid biosynthesis, yet yields only half the amount of ATP energy that the EMP pathway does. The HMP pathway also produces NADPH, which may be used for redox reactions in biosynthetic pathways. The HMP pathway does not directly produce pyruvate, however, and thus these microorganisms must also possess this portion of the EMP pathway. It is therefore not surprising that a number of microorganisms, particularly the facultative anerobes, have evolved such that they possess both of these pathways.

The ED pathway has thus far has only been found in bacteria. Although this pathway is linked partly to the HMP pathway in the reverse direction for precursor formation, the ED pathway directly forms pyruvate by the aldolase cleavage of 3-ketodeoxy-6-phosphogluconate. The ED pathway can exist on its own and is utilized by the majority of strictly aerobic microorganisms. The net result is similar to that of the HMP pathway, although one mole of ATP can be formed only if the carbon atoms are converted into pyruvate, instead of into precursor molecules.

The pyruvate molecules produced through any of these pathways can be readily converted into energy via the Krebs cycle (also known as the citric acid cycle, the citrate cycle, or the tricarboxylic acid cycle (TCA cycle)). In this process, pyruvate is first decarboxylated, resulting in the production of one molecule of NADH, 1 molecule of acetyl-CoA, and 1 molecule of CO₂. The acetyl group of acetyl CoA then reacts with

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the 4 carbon unit, oxaolacetate, leading to the formation of citric acid, a 6 carbon organic acid. Dehydration and two additional CO₂ molecules are released. Ultimately, oxaloacetate is regenerated and can serve again as an acetyl acceptor, thus completing the cycle. The electrons released during the oxidation of intermediates in the TCA cycle are transferred to NAD⁺ to yield NADH.

During respiration, the electrons from NADH are transferred to molecular oxygen or other terminal electron acceptors. This process is catalyzed by the respiratory chain, an electron transport system containing both integral membrane proteins and membrane associated proteins. This system serves two basic functions: first, to accept electrons from an electron donor and to transfer them to an electron acceptor, and second, to conserve some of the energy released during electron transfer by the synthesis of ATP. Several types of oxidation-reduction enzymes and electron transport proteins are known to be involved in such processes, including the NADH dehydrogenases, flavin-containing electron carriers, iron sulfur proteins, and cytochromes. The NADH dehydrogenases are located at the cytoplasmic surface of the plasma membrane, and transfer hydrogen atoms from NADH to flavoproteins, in turn accepting electrons from NADH. The flavoproteins are a group of electron carriers possessing a flavin prosthetic group which is alternately reduced and oxidized as it accepts and transfers electrons. Three flavins are known to participate in these reactions: riboflavin, flavin-adenine dinucleotide (FAD) and flavin-mononucleotide (FMN). Iron sulfur proteins contain a cluster of iron and sulfur atoms which are not bonded to a heme group, but which still are able to participate in dehydration and rehydration reactions. Succinate dehydrogenase and aconitase are exemplary iron-sulfur proteins; their iron-sulfur complexes serve to accept and transfer electrons as part of the overall electron-transport chain. The cytochromes are proteins containing an iron porphyrin ring (heme). There are a number of different classes of cytochromes, differing in their reduction potentials. Functionally, these cytochromes form pathways in which electrons may be transferred to other cytochromes having increasingly more positive reduction potentials. A further class of non-protein electron carriers is known: the lipid-soluble quinones (e.g., coenzyme Q). These molecules also serve as hydrogen atom acceptors and electron donors.

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The action of the respiratory chain generates a proton gradient across the cell membrane, resulting in proton motive force. This force is utilized by the cell to synthesize ATP, via the membrane-spanning enzyme, ATP synthase. This enzyme is a multiprotein complex in which the transport of H⁺ molecules through the membrane results in the physical rotation of the intracellular subunits and concomitant phosphorylation of ADP to form ATP (for review, see Fillingame, R.H. and Divall, S. (1999) *Novartis Found. Symp.* 221: 218-229, 229-234).

Non-hexose carbon substrates may also serve as carbon and energy sources for cells. Such substrates may first be converted to hexose sugars in the gluconeogenesis pathway, where glucose is first synthesized by the cell and then is degraded to produce energy. The starting material for this reaction is phosphoenolpyruvate (PEP), which is one of the key intermediates in the glycolytic pathway. PEP may be formed from substrates other than sugars, such as acetic acid, or by decarboxylation of oxaloacetate (itself an intermediate in the TCA cycle). By reversing the glycolytic pathway (utilizing a cascade of enzymes different than those of the original glycolysis pathway), glucose-6-phosphate may be formed. The conversion of pyruvate to glucose requires the utilization of 6 high energy phosphate bonds, whereas glycolysis only produces 2 ATP in the conversion of glucose to pyruvate. However, the complete oxidation of glucose (glycolysis, conversion of pyruvate into acetyl CoA, citric acid cycle, and oxidative phosphorylation) yields between 36-38 ATP, so the net loss of high energy phosphate bonds experienced during gluconeogenesis is offset by the overall greater gain in such high-energy molecules produced by the oxidation of glucose.

III. Elements and Methods of the Invention

The present invention is based, at least in part, on the discovery of novel molecules, referred to herein as SMP nucleic acid and protein molecules, which participate in the conversion of sugars to useful degradation products and energy (e.g., ATP) in C. glutamicum or which may participate in the production of useful energy-rich molecules (e.g., ATP) by other processes, such as oxidative phosphorylation. In one embodiment, the SMP molecules participate in the metabolism of carbon compounds such as sugars or the generation of energy molecules (e.g., ATP) by processes such as oxidative phosphorylation in Corynebacterium glutamicum. In a preferred embodiment,

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the activity of the SMP molecules of the present invention to contribute to carbon metabolism or energy production in C. glutamicum has an impact on the production of a desired fine chemical by this organism. In a particularly preferred embodiment, the SMP molecules of the invention are modulated in activity, such that the C. glutamicum metabolic and energetic pathways in which the SMP proteins of the invention participate are modulated in yield, production, and/or efficiency of production, which either directly or indirectly modulates the yield, production, and/or efficiency of production of a desired fine chemical by C. glutamicum.

The language, "SMP protein" or "SMP polypeptide" includes proteins which are 10 capable of performing a function involved in the metabolism of carbon compounds such as sugars and the generation of energy molecules by processes such as oxidative phosphorylation in Corynebacterium glutamicum. Examples of SMP proteins include those encoded by the SMP genes set forth in Table 1 and by the odd-numbered SEO ID NOs. The terms "SMP gene" or "SMP nucleic acid sequence" include nucleic acid 15 sequences encoding an SMP protein, which consist of a coding region and also corresponding untranslated 5' and 3' sequence regions. Examples of SMP genes include those set forth in Table 1. The terms "production" or "productivity" are art-recognized and include the concentration of the fermentation product (for example, the desired fine chemical) formed within a given time and a given fermentation volume (e.g., kg product per hour per liter). The term "efficiency of production" includes the time required for a. 20 particular level of production to be achieved (for example, how long it takes for the cell to attain a particular rate of output of a fine chemical). The term "yield" or "product/carbon yield" is art-recognized and includes the efficiency of the conversion of the carbon source into the product (i.e., fine chemical). This is generally written as, for example, kg product per kg carbon source. By increasing the yield or production of the compound, the quantity of recovered molecules, or of useful recovered molecules of that compound in a given amount of culture over a given amount of time is increased. The terms "biosynthesis" or a "biosynthetic pathway" are art-recognized and include the synthesis of a compound, preferably an organic compound, by a cell from intermediate compounds in what may be a multistep and highly regulated process. The terms "degradation" or a "degradation pathway" are art-recognized and include the breakdown of a compound, preferably an organic compound, by a cell to degradation

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products (generally speaking, smaller or less complex molecules) in what may be a multistep and highly regulated process. The term "degradation product" is art-recognized and includes breakdown products of a compound. Such products may themselves have utility as precursor (starting point) or intermediate molecules necessary for the biosynthesis of other compounds by the cell. The language "metabolism" is art-recognized and includes the totality of the biochemical reactions that take place in an organism. The metabolism of a particular compound, then, (e.g., the metabolism of an amino acid such as glycine) comprises the overall biosynthetic, modification, and degradation pathways in the cell related to this compound.

In another embodiment, the SMP molecules of the invention are capable of modulating the production of a desired molecule, such as a fine chemical, in a microorganism such as C. glutamicum. There are a number of mechanisms by which the alteration of an SMP protein of the invention may directly affect the yield, production, and/or efficiency of production of a fine chemical from a C. glutamicum strain incorporating such an altered protein. The degradation of high-energy carbon molecules such as sugars, and the conversion of compounds such as NADH and FADH₂ to more useful forms via oxidative phosphorylation results in a number of compounds which themselves may be desirable fine chemicals, such as pyruvate, ATP, NADH, and a number of intermediate sugar compounds. Further, the energy molecules (such as ATP) and the reducing equivalents (such as NADH or NADPH) produced by these metabolic pathways are utilized in the cell to drive reactions which would otherwise be energetically unfavorable. Such unfavorable reactions include many biosynthetic pathways for fine chemicals. By improving the ability of the cell to utilize a particular sugar (e.g., by manipulating the genes encoding enzymes involved in the degradation and conversion of that sugar into energy for the cell), one may increase the amount of energy available to permit unfavorable, yet desired metabolic reactions (e.g., the biosynthesis of a desired fine chemical) to occur.

The mutagenesis of one or more SMP genes of the invention may also result in SMP proteins having altered activities which indirectly impact the production of one or more desired fine chemicals from *C. glutamicum*. For example, by increasing the efficiency of utilization of one or more sugars (such that the conversion of the sugar to useful energy molecules is improved), or by increasing the efficiency of conversion of

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reducing equivalents to useful energy molecules (e.g., by improving the efficiency of oxidative phosphorylation, or the activity of the ATP synthase), one can increase the amount of these high-energy compounds available to the cell to drive normally unfavorable metabolic processes. These processes include the construction of cell walls, transcription, translation, and the biosynthesis of compounds necessary for growth and division of the cells (e.g., nucleotides, amino acids, vitamins, lipids, etc.) (Lengeler et al. (1999) Biology of Prokaryotes, Thieme Verlag: Stuttgart, p. 88-109; 913-918; 875-899). By improving the growth and multiplication of these engineered cells, it is possible to increase both the viability of the cells in large-scale culture, and also to improve their rate of division, such that a relatively larger number of cells can survive in fermentor culture. The yield, production, or efficiency of production may be increased, at least due to the presence of a greater number of viable cells, each producing the desired fine chemical. Further, a number of the degradation and intermediate compounds produced during sugar metabolism are necessary precursors and intermediates for other biosynthetic pathways throughout the cell. For example, many amino acids are synthesized directly from compounds normally resulting from glycolysis or the TCA cycle (e.g., serine is synthesized from 3-phosphoglycerate, an intermediate in glycolysis). Thus, by increasing the efficiency of conversion of sugars to useful energy molecules, it is also possible to increase the amount of useful degradation products as well.

The isolated nucleic acid sequences of the invention are contained within the genome of a *Corynebacterium glutamicum* strain available through the American Type Culture Collection, given designation ATCC 13032. The nucleotide sequence of the isolated *C. glutamicum* SMP DNAs and the predicted amino acid sequences of the *C. glutamicum* SMP proteins are shown in the Sequence Listing as odd-numbered SEQ ID NOs and even-numbered SEQ ID NOs, respectively. Computational analyses were performed which classified and/or identified these nucleotide sequences as sequences which encode proteins having a function involved in the metabolism of carbon compounds such as sugars or in the generation of energy molecules by processes such as oxidative phosphorylation in *Corynebacterium glutamicum*.

The present invention also pertains to proteins which have an amino acid sequence which is substantially homologous to an amino acid sequence of the invention

(e.g., the sequence of an even-numbered SEQ ID NO of the Sequence Listing). As used herein, a protein which has an amino acid sequence which is substantially homologous to a selected amino acid sequence is least about 50% homologous to the selected amino acid sequence, e.g., the entire selected amino acid sequence. A protein which has an amino acid sequence which is substantially homologous to a selected amino acid sequence can also be least about 50-60%, preferably at least about 60-70%, and more preferably at least about 70-80%, 80-90%, or 90-95%, and most preferably at least about 96%, 97%, 98%, 99% or more homologous to the selected amino acid sequence.

An SMP protein or a biologically active portion or fragment thereof of the invention can participate in the metabolism of carbon compounds such as sugars or in the generation of energy molecules (e.g., ATP) by processes such as oxidative phosphorylation in *Corynebacterium glutamicum*, or can have one or more of the activities set forth in Table 1.

Various aspects of the invention are described in further detail in the following subsections:

A. Isolated Nucleic Acid Molecules

One aspect of the invention pertains to isolated nucleic acid molecules that encode SMP polypeptides or biologically active portions thereof, as well as nucleic acid fragments sufficient for use as hybridization probes or primers for the identification or amplification of SMP-encoding nucleic acid (e.g., SMP DNA). As used herein, the term "nucleic acid molecule" is intended to include DNA molecules (e.g., cDNA or genomic DNA) and RNA molecules (e.g., mRNA) and analogs of the DNA or RNA generated using nucleotide analogs. This term also encompasses untranslated sequence located at both the 3' and 5' ends of the coding region of the gene: at least about 100 nucleotides of sequence upstream from the 5' end of the coding region and at least about 20 nucleotides of sequence downstream from the 3'end of the coding region of the gene. The nucleic acid molecule can be single-stranded or double-stranded, but preferably is double-stranded DNA. An "isolated" nucleic acid molecule is one which is separated from other nucleic acid molecules which are present in the natural source of the nucleic acid. Preferably, an "isolated" nucleic acid is free of sequences which naturally flank the nucleic acid (i.e., sequences located at the 5' and 3' ends of the nucleic acid) in the

genomic DNA of the organism from which the nucleic acid is derived. For example, in various embodiments, the isolated SMP nucleic acid molecule can contain less than about 5 kb, 4kb, 3kb, 2kb, 1 kb, 0.5 kb or 0.1 kb of nucleotide sequences which naturally flank the nucleic acid molecule in genomic DNA of the cell from which the nucleic acid is derived (e.g, a *C. glutamicum* cell). Moreover, an "isolated" nucleic acid molecule, such as a DNA molecule, can be substantially free of other cellular material, or culture medium when produced by recombinant techniques, or chemical precursors or other chemicals when chemically synthesized.

A nucleic acid molecule of the present invention, e.g., a nucleic acid molecule having a nucleotide sequence of an odd-numbered SEQ ID NO of the Sequence Listing, 10 or a portion thereof, can be isolated using standard molecular biology techniques and the sequence information provided herein. For example, a C. glutamicum SMP DNA can be isolated from a C. glutamicum library using all or portion of one of the odd-numbered SEQ ID NO sequences of the Sequence Listing as a hybridization probe and standard hybridization techniques (e.g., as described in Sambrook, J., Fritsh, E. F., and Maniatis, 15 T. Molecular Cloning: A Laboratory Manual. 2nd, ed., Cold Spring Harbor Laboratory, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, 1989). Moreover, a nucleic acid molecule encompassing all or a portion of one of the nucleic acid sequences of the invention (e.g., an odd-numbered SEQ ID NO:) can be isolated by the polymerase chain reaction using oligonucleotide primers designed based upon this 20 sequence (e.g., a nucleic acid molecule encompassing all or a portion of one of the nucleic acid sequences of the invention (e.g., an odd-numbered SEQ ID NO of the Sequence Listing) can be isolated by the polymerase chain reaction using oligonucleotide primers designed based upon this same sequence). For example, mRNA can be isolated from normal endothelial cells (e.g., by the guanidinium-thiocyanate 25 extraction procedure of Chirgwin et al. (1979) Biochemistry 18: 5294-5299) and DNA can be prepared using reverse transcriptase (e.g., Moloney MLV reverse transcriptase, available from Gibco/BRL, Bethesda, MD; or AMV reverse transcriptase, available from Seikagaku America, Inc., St. Petersburg, FL). Synthetic oligonucleotide primers for polymerase chain reaction amplification can be designed based upon one of the 30 nucleotide sequences shown in the Sequence Listing. A nucleic acid of the invention can be amplified using cDNA or, alternatively, genomic DNA, as a template and

WO 01/00844 PCT/IB00/00943

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- 28 -

appropriate oligonucleotide primers according to standard PCR amplification techniques. The nucleic acid so amplified can be cloned into an appropriate vector and characterized by DNA sequence analysis. Furthermore, oligonucleotides corresponding to an SMP nucleotide sequence can be prepared by standard synthetic techniques, *e.g.*, using an automated DNA synthesizer.

In a preferred embodiment, an isolated nucleic acid molecule of the invention comprises one of the nucleotide sequences shown in the Sequence Listing. The nucleic acid sequences of the invention, as set forth in the Sequence Listing, correspond to the Corynebacterium glutamicum SMP DNAs of the invention. This DNA comprises sequences encoding SMP proteins (i.e., the "coding region", indicated in each odd-numbered SEQ ID NO: sequence in the Sequence Listing), as well as 5' untranslated sequences and 3' untranslated sequences, also indicated in each odd-numbered SEQ ID NO: in the Sequence Listing. Alternatively, the nucleic acid molecule can comprise only the coding region of any of the sequences in nucleic acid sequences of the Sequence Listing.

For the purposes of this application, it will be understood that each of the nucleic acid and amino acid sequences set forth in the Sequence Listing has an identifying RXA, RXN, or RXS number having the designation "RXA," "RXN," or "RXS" followed by 5 digits (i.e., RXA01626, RXN00043, or RXS0735). Each of the nucleic acid sequences comprises up to three parts: a 5' upstream region, a coding region, and a downstream 20 region. Each of these three regions is identified by the same RXA, RXN, or RXS designation to eliminate confusion. The recitation "one of the odd-numbered sequences of the Sequence Listing", then, refers to any of the nucleic acid sequences in the Sequence Listing, which may also be distinguished by their differing RXA, RXN, or RXS designations. The coding region of each of these sequences is translated into a 25 corresponding amino acid sequence, which is also set forth in the Sequence Listing, as an even-numbered SEQ ID NO: immediately following the corresponding nucleic acid sequence. For example, the coding region for RXA02735 is set forth in SEQ ID NO:1, while the amino acid sequence which it encodes is set forth as SEQ ID NO:2. The sequences of the nucleic acid molecules of the invention are identified by the same 30 RXA, RXN, or RXS designations as the amino acid molecules which they encode, such that they can be readily correlated. For example, the amino acid sequence designated

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RXA00042 is a translation of the coding region of the nucleotide sequence of nucleic acid molecule RXA00042, and the amino acid sequence designated RXN00043 is a translation of the coding region of the nucleotide sequence of nucleic acid molecule RXN00043. The correspondence between the RXA, RXN and RXS nucleotide and amino acid sequences of the invention and their assigned SEQ ID NOs is set forth in Table 1.

Several of the genes of the invention are "F-designated genes". An F-designated gene includes those genes set forth in Table 1 which have an 'F' in front of the RXAdesignation. For example, SEQ ID NO:11, designated, as indicated on Table 1, as "F RXA01312", is an F-designated gene, as are SEQ ID NOs: 29, 33, and 39 (designated on Table 1 as "F RXA02803", "F RXA02854", and "F RXA01365", respectively).

In one embodiment, the nucleic acid molecules of the present invention are not intended to include those compiled in Table 2. In the case of the dapD gene, a sequence for this gene was published in Wehrmann, A., et al. (1998) J. Bacteriol. 180(12): 3159-3165. However, the sequence obtained by the inventors of the present application is significantly longer than the published version. It is believed that the published version relied on an incorrect start codon, and thus represents only a fragment of the actual coding region.

In another preferred embodiment, an isolated nucleic acid molecule of the invention comprises a nucleic acid molecule which is a complement of one of the nucleotide sequences of the invention (e.g., a sequence of an odd-numbered SEQ ID NO: of the Sequence Listing), or a portion thereof. A nucleic acid molecule which is complementary to one of the nucleotide sequences of the invention is one which is sufficiently complementary to one of the nucleotide sequences shown in the Sequence Listing (e.g., the sequence of an odd-numbered SEQ ID NO:) such that it can hybridize to one of the nucleotide sequences of the invention, thereby forming a stable duplex.

In still another preferred embodiment, an isolated nucleic acid molecule of the invention comprises a nucleotide sequence which is at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, or 60%, preferably at least about 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, or 70%, more preferably at least about 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, or 80%, 81%, 82%, 83%, 84%, 85%, 86%,

PCT/IB00/00943 WO 01/00844

- 30 -

87%, 88%, 89%, or 90%, or 91%, 92%, 93%, 94%, and even more preferably at least about 95%, 96%, 97%, 98%, 99% or more homologous to a nucleotide sequence of the invention (e.g., a sequence of an odd-numbered SEQ ID NO: of the Sequence Listing), or a portion thereof. Ranges and identity values intermediate to the above-recited ranges, (e.g., 70-90% identical or 80-95% identical) are also intended to be encompassed by the present invention. For example, ranges of identity values using a combination of any of the above values recited as upper and/or lower limits are intendedto be included. In an additional preferred embodiment, an isolated nucleic acid molecule of the invention comprises a nucleotide sequence which hybridizes, e.g., hybridizes under stringent conditions, to one of the nucleotide sequences of the invention, or a portion thereof.

Moreover, the nucleic acid molecule of the invention can comprise only a portion of the coding region of the sequence of one of the odd-numbered SEQ ID NOs of the Sequence Listing, for example a fragment which can be used as a probe or primer 15 or a fragment encoding a biologically active portion of an SMP protein. The nucleotide sequences determined from the cloning of the SMP genes from C. glutamicum allows for the generation of probes and primers designed for use in identifying and/or cloning SMP homologues in other cell types and organisms, as well as SMP homologues from other Corynebacteria or related species. The probe/primer typically comprises substantially purified oligonucleotide. The oligonucleotide typically comprises a region of nucleotide sequence that hybridizes under stringent conditions to at least about 12, preferably about 25, more preferably about 40, 50 or 75 consecutive nucleotides of a sense strand of one of the nucleotide sequences of the invention (e.g., a sequence of one of the odd-numbered SEQ ID NOs of the Sequence Listing), an anti-sense sequence of one of these sequences, or naturally occurring mutants thereof. Primers based on a nucleotide sequence of the invention can be used in PCR reactions to clone SMP homologues. Probes based on the SMP nucleotide sequences can be used to detect transcripts or genomic sequences encoding the same or homologous proteins. In preferred embodiments, the probe further comprises a label group attached thereto, e.g. the label group can be a radioisotope, a fluorescent compound, an enzyme, or an enzyme co-factor. Such probes can be used as a part of a diagnostic test kit for identifying cells which misexpress an SMP protein, such as by measuring a level of an SMP-encoding

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nucleic acid in a sample of cells, e.g., detecting SMP mRNA levels or determining whether a genomic SMP gene has been mutated or deleted.

In one embodiment, the nucleic acid molecule of the invention encodes a protein or portion thereof which includes an amino acid sequence which is sufficiently homologous to an amino acid sequence of the invention (e.g., a sequence of an evennumbered SEQ ID NO of the Sequence Listing) such that the protein or portion thereof maintains the ability to perform a function involved in the metabolism of carbon compounds such as sugars or in the generation of energy molecules (e.g., ATP) by processes such as oxidative phosphorylation in Corynebacterium glutamicum. As used herein, the language "sufficiently homologous" refers to proteins or portions thereof which have amino acid sequences which include a minimum number of identical or equivalent (e.g., an amino acid residue which has a similar side chain as an amino acid residue in a sequence of one of the even-numbered SEQ ID NOs of the Sequence Listing) amino acid residues to an amino acid sequence of the invention such that the protein or portion thereof is able to perform a function involved in the metabolism of 15 carbon compounds such as sugars or in the generation of energy molecules (e.g., ATP) by processes such as oxidative phosphorylation in Corynebacterium glutamicum. Protein members of such sugar metabolic pathways or energy producing systems, as described herein, may play a role in the production and secretion of one or more fine chemicals. Examples of such activities are also described herein. Thus, "the function of 20 an SMP protein" contributes either directly or indirectly to the yield, production, and/or efficiency of production of one or more fine chemicals. Examples of SMP protein activities are set forth in Table 1.

In another embodiment, the protein is at least about 50-60%, preferably at least about 60-70%, and more preferably at least about 70-80%, 80-90%, 90-95%, and most preferably at least about 96%, 97%, 98%, 99% or more homologous to an entire amino acid sequence of the invention(e.g., a sequence of an even-numbered SEQ ID NO: of the Sequence Listing).

Portions of proteins encoded by the SMP nucleic acid molecules of the invention are preferably biologically active portions of one of the SMP proteins. As used herein, the term "biologically active portion of an SMP protein" is intended to include a portion, e.g., a domain/motif, of an SMP protein that participates in the metabolism of carbon

WO 01/00844 PCT/IB00/00943

- 32 -

compounds such as sugars, or in energy-generating pathways in *C. glutamicum*, or has an activity as set forth in Table 1. To determine whether an SMP protein or a biologically active portion thereof can participate in the metabolism of carbon compounds or in the production of energy-rich molecules in *C. glutamicum*, an assay of enzymatic activity may be performed. Such assay methods are well known to those of ordinary skill in the art, as detailed in Example 8 of the Exemplification.

Additional nucleic acid fragments encoding biologically active portions of an SMP protein can be prepared by isolating a portion of one of the amino acid sequences of the invention (e.g., a sequence of an even-numbered SEQ ID NO: of the Sequence Listing), expressing the encoded portion of the SMP protein or peptide (e.g., by recombinant expression in vitro) and assessing the activity of the encoded portion of the SMP protein or peptide.

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The invention further encompasses nucleic acid molecules that differ from one of the nucleotide sequences of the invention (e.g., a sequence of an odd-numbered SEQ ID NO: of the Sequence Listing) (and portions thereof) due to degeneracy of the genetic code and thus encode the same SMP protein as that encoded by the nucleotide sequences of the invention. In another embodiment, an isolated nucleic acid molecule of the invention has a nucleotide sequence encoding a protein having an amino acid sequence shown in the Sequence Listing (e.g., an even-numbered SEQ ID NO:). In a still further embodiment, the nucleic acid molecule of the invention encodes a full length C. glutamicum protein which is substantially homologous to an amino acid of the invention (encoded by an open reading frame shown in an odd-numbered SEQ ID NO: of the Sequence Listing).

It will be understood by one of ordinary skill in the art that in one embodiment the sequences of the invention are not meant to include the sequences of the prior art, such as those Genbank sequences set forth in Tables 2 or 4 which were available prior to the present invention. In one embodiment, the invention includes nucleotide and amino acid sequences having a percent identity to a nucleotide or amino acid sequence of the invention which is greater than that of a sequence of the prior art (e.g., a Genbank sequence (or the protein encoded by such a sequence) set forth in Tables 2 or 4). For example, the invention includes a nucleotide sequence which is greater than and/or at least 58% identical to the nucleotide sequence designated RXA00014 (SEQ ID NO:41),

a nucleotide sequence which is greater than and/or at least % identical to the nucleotide sequence designated RXA00195 (SEQ ID NO:399), and a nucleotide sequence which is greater than and/or at least 42% identical to the nucleotide sequence designated RXA00196 (SEQ ID NO:401). One of ordinary skill in the art would be able to calculate the lower threshold of percent identity for any given sequence of the invention by examining the GAP-calculated percent identity scores set forth in Table 4 for each of the three top hits for the given sequence, and by subtracting the highest GAP-calculated percent identity from 100 percent. One of ordinary skill in the art will also appreciate that nucleic acid and amino acid sequences having percent identities greater than the 10 lower threshold so calculated (e.g., at least 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, or 60%, preferably at least about 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, or 70%, more preferably at least about 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, or 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, or 90%, or 91%, 92%, 93%, 94%, and even more preferably at least about 95%, 96%, 97%, 15 98%, 99% or more identical) are also encompassed by the invention.

In addition to the *C. glutamicum* SMP nucleotide sequences set forth in the Sequence Listing as odd-numbered SEQ ID NOs, it will be appreciated by those of ordinary skill in the art that DNA sequence polymorphisms that lead to changes in the amino acid sequences of SMP proteins may exist within a population (*e.g.*, the *C. glutamicum* population). Such genetic polymorphism in the SMP gene may exist among individuals within a population due to natural variation. As used herein, the terms "gene" and "recombinant gene" refer to nucleic acid molecules comprising an open reading frame encoding an SMP protein, preferably a *C. glutamicum* SMP protein. Such natural variations can typically result in 1-5% variance in the nucleotide sequence of the SMP gene. Any and all such nucleotide variations and resulting amino acid polymorphisms in SMP that are the result of natural variation and that do not alter the functional activity of SMP proteins are intended to be within the scope of the invention.

Nucleic acid molecules corresponding to natural variants and non-C. glutamicum. homologues of the C. glutamicum SMP DNA of the invention can be isolated based on their homology to the C. glutamicum SMP nucleic acid disclosed herein using the C. glutamicum DNA, or a portion thereof, as a hybridization probe according to standard hybridization techniques under stringent hybridization conditions. Accordingly, in

WO 01/00844 PCT/IB00/00943

- 34 -

another embodiment, an isolated nucleic acid molecule of the invention is at least 15 nucleotides in length and hybridizes under stringent conditions to the nucleic acid molecule comprising a nucleotide sequence of of an odd-numbered SEQ ID NO: of the Sequence Listing. In other embodiments, the nucleic acid is at least 30, 50, 100, 250 or more nucleotides in length. As used herein, the term "hybridizes under stringent conditions" is intended to describe conditions for hybridization and washing under which nucleotide sequences at least 60% homologous to each other typically remain hybridized to each other. Preferably, the conditions are such that sequences at least about 65%, more preferably at least about 70%, and even more preferably at least about 75% or more homologous to each other typically remain hybridized to each other. Such stringent conditions are known to those of ordinary skill in the art and can be found in Current Protocols in Molecular Biology, John Wiley & Sons, N.Y. (1989), 6.3.1-6.3.6. A preferred, non-limiting example of stringent hybridization conditions are hybridization in 6X sodium chloride/sodium citrate (SSC) at about 45°C, followed by one or more washes in 0.2 X SSC, 0.1% SDS at 50-65°C. Preferably, an isolated nucleic acid molecule of the invention that hybridizes under stringent conditions to a nucleotide sequence of the invention corresponds to a naturally-occurring nucleic acid molecule. As used herein, a "naturally-occurring" nucleic acid molecule refers to an RNA or DNA molecule having a nucleotide sequence that occurs in nature (e.g., encodes a natural protein). In one embodiment, the nucleic acid encodes a natural C. glutamicum SMP protein.

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In addition to naturally-occurring variants of the SMP sequence that may exist in the population, one of ordinary skill in the art will further appreciate that changes can be introduced by mutation into a nucleotide sequence of the invention, thereby leading to changes in the amino acid sequence of the encoded SMP protein, without altering the functional ability of the SMP protein. For example, nucleotide substitutions leading to amino acid substitutions at "non-essential" amino acid residues can be made in a nucleotide sequence of the invention. A "non-essential" amino acid residue is a residue, that can be altered from the wild-type sequence of one of the SMP proteins (e.g., an even-numbered SEQ ID NO: of the Sequence Listing) without altering the activity of said SMP protein, whereas an "essential" amino acid residue is required for SMP protein activity. Other amino acid residues, however, (e.g., those that are not conserved or only

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semi-conserved in the domain having SMP activity) may not be essential for activity and thus are likely to be amenable to alteration without altering SMP activity.

Accordingly, another aspect of the invention pertains to nucleic acid molecules encoding SMP proteins that contain changes in amino acid residues that are not essential for SMP activity. Such SMP proteins differ in amino acid sequence from a sequence of an even-numbered SEQ ID NO: of the Sequence Listing yet retain at least one of the SMP activities described herein. In one embodiment, the isolated nucleic acid molecule comprises a nucleotide sequence encoding a protein, wherein the protein comprises an amino acid sequence at least about 50% homologous to an amino acid sequence of the invention and is capable of participate in the metabolism of carbon compounds such as sugars, or in the biosynthesis of high-energy compounds in C. glutamicum, or has one or more activities set forth in Table 1. Preferably, the protein encoded by the nucleic acid molecule is at least about 50-60% homologous to the amino acid sequence of one of the odd-numbered SEQ ID NOs of the Sequence Listing, more preferably at least about 60-70% homologous to one of these sequences, even more preferably at least about 70-80%, 80-90%, 90-95% homologous to one of these sequences, and most preferably at least about 96%, 97%, 98%, or 99% homologous to one of the amino acid sequences of the invention.

To determine the percent homology of two amino acid sequences (e.g., one of the amino acid sequences of the invention and a mutant form thereof) or of two nucleic 20 acids, the sequences are aligned for optimal comparison purposes (e.g., gaps can be introduced in the sequence of one protein or nucleic acid for optimal alignment with the other protein or nucleic acid). The amino acid residues or nucleotides at corresponding amino acid positions or nucleotide positions are then compared. When a position in one sequence (e.g., one of the amino acid sequences the invention) is occupied by the same amino acid residue or nucleotide as the corresponding position in the other sequence (e.g., a mutant form of the amino acid sequence), then the molecules are homologous at that position (i.e., as used herein amino acid or nucleic acid "homology" is equivalent to amino acid or nucleic acid "identity"). The percent homology between the two sequences is a function of the number of identical positions shared by the sequences (i.e., % homology = # of identical positions/total # of positions x 100).

An isolated nucleic acid molecule encoding an SMP protein homologous to a protein sequence of the invention (e.g., a sequence of an even-numbered SEQ ID NO: of the Sequence Listing) can be created by introducing one or more nucleotide substitutions, additions or deletions into a nucleotide sequence of the invention such that one or more amino acid substitutions, additions or deletions are introduced into the encoded protein. Mutations can be introduced into one of the nucleotide sequences of the invention by standard techniques, such as site-directed mutagenesis and PCRmediated mutagenesis. Preferably, conservative amino acid substitutions are made at one or more predicted non-essential amino acid residues. A "conservative amino acid substitution" is one in which the amino acid residue is replaced with an amino acid residue having a similar side chain. Families of amino acid residues having similar side chains have been defined in the art. These families include amino acids with basic side chains (e.g., lysine, arginine, histidine), acidic side chains (e.g., aspartic acid, glutamic acid), uncharged polar side chains (e.g., glycine, asparagine, glutamine, serine, threonine, tyrosine, cysteine), nonpolar side chains (e.g., alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, tryptophan), beta-branched side chains (e.g., threonine, valine, isoleucine) and aromatic side chains (e.g., tyrosine, phenylalanine, tryptophan, histidine). Thus, a predicted nonessential amino acid residue in an SMP protein is preferably replaced with another amino acid residue from the same side chain family. Alternatively, in another embodiment, mutations can be introduced randomly along all or part of an SMP coding sequence, such as by saturation mutagenesis, and the resultant mutants can be screened for an SMP activity described herein to identify mutants that retain SMP activity. Following mutagenesis of the nucleotide sequence of one of the odd-numbered SEQ ID NOs of the Sequence Listing, the encoded protein can be expressed recombinantly and the activity of the protein can be determined using, for example, assays described herein (see Example 8 of the Exemplification).

In addition to the nucleic acid molecules encoding SMP proteins described above, another aspect of the invention pertains to isolated nucleic acid molecules which are antisense thereto. An "antisense" nucleic acid comprises a nucleotide sequence which is complementary to a "sense" nucleic acid encoding a protein, e.g., complementary to the coding strand of a double-stranded DNA molecule or

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complementary to an mRNA sequence. Accordingly, an antisense nucleic acid can hydrogen bond to a sense nucleic acid. The antisense nucleic acid can be complementary to an entire SMP coding strand, or to only a portion thereof. In one embodiment, an antisense nucleic acid molecule is antisense to a "coding region" of the coding strand of a nucleotide sequence encoding an SMP protein. The term "coding region" refers to the region of the nucleotide sequence comprising codons which are translated into amino acid residues (e.g., the entire coding region of NO. 3 (RXA01626) comprises nucleotides 1 to 345). In another embodiment, the antisense nucleic acid molecule is antisense to a "noncoding region" of the coding strand of a nucleotide sequence encoding SMP. The term "noncoding region" refers to 5' and 3' sequences which flank the coding region that are not translated into amino acids (i.e., also referred to as 5' and 3' untranslated regions).

Given the coding strand sequences encoding SMP disclosed herein (e.g., the sequences set forth as odd-numbered SEQ ID NOs in the Sequence Listing), antisense nucleic acids of the invention can be designed according to the rules of Watson and 15 Crick base pairing. The antisense nucleic acid molecule can be complementary to the entire coding region of SMP mRNA, but more preferably is an oligonucleotide which is antisense to only a portion of the coding or noncoding region of SMP mRNA. For example, the antisense oligonucleotide can be complementary to the region surrounding the translation start site of SMP mRNA. An antisense oligonucleotide can be, for 20 example, about 5, 10, 15, 20, 25, 30, 35, 40, 45 or 50 nucleotides in length. An antisense nucleic acid of the invention can be constructed using chemical synthesis and enzymatic ligation reactions using procedures known in the art. For example, an antisense nucleic acid (e.g., an antisense oligonucleotide) can be chemically synthesized using naturally occurring nucleotides or variously modified nucleotides designed to increase the biological stability of the molecules or to increase the physical stability of the duplex formed between the antisense and sense nucleic acids, e.g., phosphorothioate derivatives and acridine substituted nucleotides can be used. Examples of modified nucleotides which can be used to generate the antisense nucleic acid include 5fluorouracil, 5-bromouracil, 5-iodouracil, hypoxanthine, xanthine, 4acetylcytosine, 5-(carboxyhydroxylmethyl) uracil, 5-carboxymethylaminomethyl-2thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-

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galactosylqueosine, inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-

methoxycarboxymethyluracil, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil, 3-(3-amino-3-N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine. Alternatively, the antisense nucleic acid can be produced biologically using an expression vector into which a nucleic acid has been subcloned in an antisense orientation (*i.e.*, RNA transcribed from the inserted nucleic acid will be of an antisense orientation to a target nucleic acid of interest, described further in the following subsection).

The antisense nucleic acid molecules of the invention are typically administered to a cell or generated *in situ* such that they hybridize with or bind to cellular mRNA and/or genomic DNA encoding an SMP protein to thereby inhibit expression of the protein, *e.g.*, by inhibiting transcription and/or translation. The hybridization can be by conventional nucleotide complementarity to form a stable duplex, or, for example, in the case of an antisense nucleic acid molecule which binds to DNA duplexes, through specific interactions in the major groove of the double helix. The antisense molecule can be modified such that it specifically binds to a receptor or an antigen expressed on a selected cell surface, *e.g.*, by linking the antisense nucleic acid molecule to a peptide or an antibody which binds to a cell surface receptor or antigen. The antisense nucleic acid molecule can also be delivered to cells using the vectors described herein. To achieve sufficient intracellular concentrations of the antisense molecules, vector constructs in which the antisense nucleic acid molecule is placed under the control of a strong prokaryotic, viral, or eukaryotic promoter are preferred.

In yet another embodiment, the antisense nucleic acid molecule of the invention is an α -anomeric nucleic acid molecule. An α -anomeric nucleic acid molecule forms specific double-stranded hybrids with complementary RNA in which, contrary to the usual β -units, the strands run parallel to each other (Gaultier *et al.* (1987) *Nucleic Acids*. *Res.* 15:6625-6641). The antisense nucleic acid molecule can also comprise a 2'-o-

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methylribonucleotide (Inoue et al. (1987) Nucleic Acids Res. 15:6131-6148) or a chimeric RNA-DNA analogue (Inoue et al. (1987) FEBS Lett. 215:327-330).

In still another embodiment, an antisense nucleic acid of the invention is a ribozyme. Ribozymes are catalytic RNA molecules with ribonuclease activity which are capable of cleaving a single-stranded nucleic acid, such as an mRNA, to which they have a complementary region. Thus, ribozymes (e.g., hammerhead ribozymes (described in Haselhoff and Gerlach (1988) Nature 334:585-591)) can be used to catalytically cleave SMP mRNA transcripts to thereby inhibit translation of SMP mRNA. A ribozyme having specificity for an SMP-encoding nucleic acid can be designed based upon the nucleotide sequence of an SMP cDNA disclosed herein (i.e., SEQ ID NO. 3 (RXA01626)). For example, a derivative of a Tetrahymena L-19 IVS RNA can be constructed in which the nucleotide sequence of the active site is complementary to the nucleotide sequence to be cleaved in an SMP-encoding mRNA. See, e.g., Cech et al. U.S. Patent No. 4,987,071 and Cech et al. U.S. Patent No. 5,116,742. Alternatively, SMP mRNA can be used to select a catalytic RNA having a specific ribonuclease activity from a pool of RNA molecules. See, e.g., Bartel, D. and Szostak, J.W. (1993) Science 261:1411-1418.

Alternatively, SMP gene expression can be inhibited by targeting nucleotide sequences complementary to the regulatory region of an SMP nucleotide sequence (e.g., an SMP promoter and/or enhancers) to form triple helical structures that prevent transcription of an SMP gene in target cells. See generally, Helene, C. (1991)

Anticancer Drug Des. 6(6):569-84; Helene, C. et al. (1992) Ann. N.Y. Acad. Sci. 660:27-36; and Maher, L.J. (1992) Bioassays 14(12):807-15.

25 B. Recombinant Expression Vectors and Host Cells

Another aspect of the invention pertains to vectors, preferably expression vectors, containing a nucleic acid encoding an SMP protein (or a portion thereof). As used herein, the term "vector" refers to a nucleic acid molecule capable of transporting another nucleic acid to which it has been linked. One type of vector is a "plasmid", which refers to a circular double stranded DNA loop into which additional DNA segments can be ligated. Another type of vector is a viral vector, wherein additional DNA segments can be ligated into the viral genome. Certain vectors are capable of

PCT/IB00/00943 WO 01/00844

- 40 -

autonomous replication in a host cell into which they are introduced (e.g., bacterial vectors having a bacterial origin of replication and episomal mammalian vectors). Other vectors (e.g., non-episomal mammalian vectors) are integrated into the genome of a host cell upon introduction into the host cell, and thereby are replicated along with the host genome. Moreover, certain vectors are capable of directing the expression of genes to which they are operatively linked. Such vectors are referred to herein as "expression vectors". In general, expression vectors of utility in recombinant DNA techniques are often in the form of plasmids. In the present specification, "plasmid" and "vector" can be used interchangeably as the plasmid is the most commonly used form of vector. However, the invention is intended to include such other forms of expression vectors,

such as viral vectors (e.g., replication defective retroviruses, adenoviruses and adenoassociated viruses), which serve equivalent functions.

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The recombinant expression vectors of the invention comprise a nucleic acid of the invention in a form suitable for expression of the nucleic acid in a host cell, which means that the recombinant expression vectors include one or more regulatory sequences, selected on the basis of the host cells to be used for expression, which is operatively linked to the nucleic acid sequence to be expressed. Within a recombinant expression vector, "operably linked" is intended to mean that the nucleotide sequence of interest is linked to the regulatory sequence(s) in a manner which allows for expression of the nucleotide sequence (e.g., in an in vitro transcription/translation system or in a host cell when the vector is introduced into the host cell). The term "regulatory sequence" is intended to include promoters, enhancers and other expression control elements (e.g., polyadenylation signals). Such regulatory sequences are described, for example, in Goeddel; Gene Expression Technology: Methods in Enzymology 185, Academic Press, San Diego, CA (1990). Regulatory sequences include those which direct constitutive expression of a nucleotide sequence in many types of host cell and those which direct expression of the nucleotide sequence only in certain host cells. Preferred regulatory sequences are, for example, promoters such as cos-, tac-, trp-, tet-, trp-tet-, lpp-, lac-, lpp-lac-, lacI^q-, T7-, T5-, T3-, gal-, trc-, ara-, SP6-, arny, SPO2, λ-P_Ror λ P₁, which are used preferably in bacteria. Additional regulatory sequences are, for example, promoters from yeasts and fungi, such as ADC1, MFa, AC, P-60, CYC1, GAPDH, TEF, rp28, ADH, promoters from plants such as CaMV/35S, SSU, OCS, lib4,

WO 01/00844 PCT/IB00/00943

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usp, STLS1, B33, nos or ubiquitin- or phaseolin-promoters. It is also possible to use artificial promoters. It will be appreciated by those of ordinary skill in the art that the design of the expression vector can depend on such factors as the choice of the host cell to be transformed, the level of expression of protein desired, etc. The expression vectors of the invention can be introduced into host cells to thereby produce proteins or peptides, including fusion proteins or peptides, encoded by nucleic acids as described herein (e.g., SMP proteins, mutant forms of SMP proteins, fusion proteins, etc.).

The recombinant expression vectors of the invention can be designed for expression of SMP proteins in prokaryotic or eukaryotic cells. For example, SMP genes can be expressed in bacterial cells such as C. glutamicum, insect cells (using baculovirus expression vectors), yeast and other fungal cells (see Romanos, M.A. et al. (1992) "Foreign gene expression in yeast: a review", Yeast 8: 423-488; van den Hondel, C.A.M.J.J. et al. (1991) "Heterologous gene expression in filamentous fungi" in: More Gene Manipulations in Fungi, J.W. Bennet & L.L. Lasure, eds., p. 396-428: Academic Press: San Diego; and van den Hondel, C.A.M.J.J. & Punt, P.J. (1991) "Gene transfer systems and vector development for filamentous fungi, in: Applied Molecular Genetics of Fungi, Peberdy, J.F. et al., eds., p. 1-28, Cambridge University Press: Cambridge), algae and multicellular plant cells (see Schmidt, R. and Willmitzer, L. (1988) High efficiency Agrobacterium tumefaciens - mediated transformation of Arabidopsis thaliana leaf and cotyledon explants" Plant Cell Rep: 583-586), or mammalian cells. Suitable host cells are discussed further in Goeddel, Gene Expression Technology: Methods in Enzymology 185, Academic Press, San Diego, CA (1990). Alternatively, the recombinant expression vector can be transcribed and translated in vitro, for example using T7 promoter regulatory sequences and T7 polymerase.

Expression of proteins in prokaryotes is most often carried out with vectors containing constitutive or inducible promoters directing the expression of either fusion or non-fusion proteins. Fusion vectors add a number of amino acids to a protein encoded therein, usually to the amino terminus of the recombinant protein but also to the C-terminus or fused within suitable regions in the proteins. Such fusion vectors typically serve three purposes: 1) to increase expression of recombinant protein; 2) to increase the solubility of the recombinant protein; and 3) to aid in the purification of the recombinant protein by acting as a ligand in affinity purification. Often, in fusion

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expression vectors, a proteolytic cleavage site is introduced at the junction of the fusion moiety and the recombinant protein to enable separation of the recombinant protein from the fusion moiety subsequent to purification of the fusion protein. Such enzymes, and their cognate recognition sequences, include Factor Xa, thrombin and enterokinase.

Typical fusion expression vectors include pGEX (Pharmacia Biotech Inc; Smith, D.B. and Johnson, K.S. (1988) *Gene* 67:31-40), pMAL (New England Biolabs, Beverly, MA) and pRIT5 (Pharmacia, Piscataway, NJ) which fuse glutathione S-transferase (GST), maltose E binding protein, or protein A, respectively, to the target recombinant protein. In one embodiment, the coding sequence of the SMP protein is cloned into a pGEX expression vector to create a vector encoding a fusion protein comprising, from the N-terminus to the C-terminus, GST-thrombin cleavage site-X protein. The fusion protein can be purified by affinity chromatography using glutathione-agarose resin. Recombinant SMP protein unfused to GST can be recovered by cleavage of the fusion protein with thrombin.

Examples of suitable inducible non-fusion E. coli expression vectors include pTrc (Amann et al., (1988) Gene 69:301-315), pLG338, pACYC184, pBR322, pUC18, pUC19, pKC30, pRep4, pHS1, pHS2, pPLc236, pMBL24, pLG200, pUR290, pIN-III113-B1, \(\lambda\)gt11, pBdCl, and pET 11d (Studier et al., Gene Expression Technology: Methods in Enzymology 185, Academic Press, San Diego, California (1990) 60-89; and Pouwels et al., eds. (1985) Cloning Vectors. Elsevier: New York IBSN 0 444 904018). Target gene expression from the pTrc vector relies on host RNA polymerase transcription from a hybrid trp-lac fusion promoter. Target gene expression from the pET 11d vector relies on transcription from a T7 gn10-lac fusion promoter mediated by a coexpressed viral RNA polymerase (T7 gn1). This viral polymerase is supplied by host strains BL21(DE3) or HMS174(DE3) from a resident λ prophage harboring a T7 gn1 gene under the transcriptional control of the lacUV 5 promoter. For transformation of other varieties of bacteria, appropriate vectors may be selected. For example, the plasmids pIJ101, pIJ364, pIJ702 and pIJ361 are known to be useful in transforming Streptomyces, while plasmids pUB110, pC194, or pBD214 are suited for transformation of Bacillus species. Several plasmids of use in the transfer of genetic information into Corynebacterium include pHM1519, pBL1, pSA77, or pAJ667 (Pouwels et al., eds. (1985) Cloning Vectors. Elsevier: New York IBSN 0 444 904018).

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One strategy to maximize recombinant protein expression is to express the protein in a host bacteria with an impaired capacity to proteolytically cleave the recombinant protein (Gottesman, S., Gene Expression Technology: Methods in Enzymology 185, Academic Press, San Diego, California (1990) 119-128). Another strategy is to alter the nucleic acid sequence of the nucleic acid to be inserted into an expression vector so that the individual codons for each amino acid are those preferentially utilized in the bacterium chosen for expression, such as C. glutamicum (Wada et al. (1992) Nucleic Acids Res. 20:2111-2118). Such alteration of nucleic acid sequences of the invention can be carried out by standard DNA synthesis techniques.

In another embodiment, the SMP protein expression vector is a yeast expression vector. Examples of vectors for expression in yeast *S. cerevisiae* include pYepSec1 (Baldari, *et al.*, (1987) *Embo J.* 6:229-234), 2 μ, pAG-1, Yep6, Yep13, pEMBLYe23, pMFa (Kurjan and Herskowitz, (1982) *Cell* 30:933-943), pJRY88 (Schultz *et al.*, (1987) *Gene* 54:113-123), and pYES2 (Invitrogen Corporation, San Diego, CA). Vectors and methods for the construction of vectors appropriate for use in other fungi, such as the filamentous fungi, include those detailed in: van den Hondel, C.A.M.J.J. & Punt, P.J. (1991) "Gene transfer systems and vector development for filamentous fungi, in: Applied Molecular Genetics of Fungi, J.F. Peberdy, *et al.*, eds., p. 1-28, Cambridge University Press: Cambridge, and Pouwels *et al.*, eds. (1985) Cloning Vectors. Elsevier: New York (IBSN 0 444 904018).

Alternatively, the SMP proteins of the invention can be expressed in insect cells using baculovirus expression vectors. Baculovirus vectors available for expression of proteins in cultured insect cells (e.g., Sf 9 cells) include the pAc series (Smith et al. (1983) Mol. Cell Biol. 3:2156-2165) and the pVL series (Lucklow and Summers (1989) Virology 170:31-39).

In another embodiment, the SMP proteins of the invention may be expressed in unicellular plant cells (such as algae) or in plant cells from higher plants (e.g., the spermatophytes, such as crop plants). Examples of plant expression vectors include those detailed in: Becker, D., Kemper, E., Schell, J. and Masterson, R. (1992) "New plant binary vectors with selectable markers located proximal to the left border", *Plant Mol. Biol.* 20: 1195-1197; and Bevan, M.W. (1984) "Binary *Agrobacterium* vectors for plant transformation", *Nucl. Acid. Res.* 12: 8711-8721, and include pLGV23, pGHlac+,

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pBIN19, pAK2004, and pDH51 (Pouwels et al., eds. (1985) Cloning Vectors. Elsevier: New York IBSN 0 444 904018).

In yet another embodiment, a nucleic acid of the invention is expressed in mammalian cells using a mammalian expression vector. Examples of mammalian expression vectors include pCDM8 (Seed, B. (1987) *Nature* 329:840) and pMT2PC (Kaufman *et al.* (1987) *EMBO J.* 6:187-195). When used in mammalian cells, the expression vector's control functions are often provided by viral regulatory elements. For example, commonly used promoters are derived from polyoma, Adenovirus 2, cytomegalovirus and Simian Virus 40. For other suitable expression systems for both prokaryotic and eukaryotic cells see chapters 16 and 17 of Sambrook, J., Fritsh, E. F., and Maniatis, T. *Molecular Cloning: A Laboratory Manual. 2nd, ed., Cold Spring Harbor Laboratory*, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, 1989.

In another embodiment, the recombinant mammalian expression vector is capable of directing expression of the nucleic acid preferentially in a particular cell type 15 (e.g., tissue-specific regulatory elements are used to express the nucleic acid). Tissuespecific regulatory elements are known in the art. Non-limiting examples of suitable tissue-specific promoters include the albumin promoter (liver-specific; Pinkert et al. (1987) Genes Dev. 1:268-277), lymphoid-specific promoters (Calame and Eaton (1988) Adv. Immunol. 43:235-275), in particular promoters of T cell receptors (Winoto and 20 Baltimore (1989) EMBO J. 8:729-733) and immunoglobulins (Banerji et al. (1983) Cell 33:729-740; Queen and Baltimore (1983) Cell 33:741-748), neuron-specific promoters (e.g., the neurofilament promoter; Byrne and Ruddle (1989) PNAS 86:5473-5477), pancreas-specific promoters (Edlund et al. (1985) Science 230:912-916), and mammary 25 gland-specific promoters (e.g., milk whey promoter; U.S. Patent No. 4,873,316 and European Application Publication No. 264,166). Developmentally-regulated promoters are also encompassed, for example the murine hox promoters (Kessel and Gruss (1990) Science 249:374-379) and the α-fetoprotein promoter (Campes and Tilghman (1989) Genes Dev. 3:537-546).

The invention further provides a recombinant expression vector comprising a DNA molecule of the invention cloned into the expression vector in an antisense orientation. That is, the DNA molecule is operatively linked to a regulatory sequence in

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a manner which allows for expression (by transcription of the DNA molecule) of an RNA molecule which is antisense to SMP mRNA. Regulatory sequences operatively linked to a nucleic acid cloned in the antisense orientation can be chosen which direct the continuous expression of the antisense RNA molecule in a variety of cell types, for instance viral promoters and/or enhancers, or regulatory sequences can be chosen which direct constitutive, tissue specific or cell type specific expression of antisense RNA. The antisense expression vector can be in the form of a recombinant plasmid, phagemid or attenuated virus in which antisense nucleic acids are produced under the control of a high efficiency regulatory region, the activity of which can be determined by the cell type into which the vector is introduced. For a discussion of the regulation of gene expression using antisense genes see Weintraub, H. et al., Antisense RNA as a molecular tool for genetic analysis, Reviews - Trends in Genetics, Vol. 1(1) 1986.

Another aspect of the invention pertains to host cells into which a recombinant expression vector of the invention has been introduced. The terms "host cell" and "recombinant host cell" are used interchangeably herein. It is understood that such terms refer not only to the particular subject cell but to the progeny or potential progeny of such a cell. Because certain modifications may occur in succeeding generations due to either mutation or environmental influences, such progeny may not, in fact, be identical to the parent cell, but are still included within the scope of the term as used herein.

A host cell can be any prokaryotic or eukaryotic cell. For example, an SMP protein can be expressed in bacterial cells such as *C. glutamicum*, insect cells, yeast or mammalian cells (such as Chinese hamster ovary cells (CHO) or COS cells). Other suitable host cells are known to one of ordinary skill in the art. Microorganisms related to *Corynebacterium glutamicum* which may be conveniently used as host cells for the nucleic acid and protein molecules of the invention are set forth in Table 3.

Vector DNA can be introduced into prokaryotic or eukaryotic cells via conventional transformation or transfection techniques. As used herein, the terms "transformation" and "transfection", "conjugation" and "transduction" are intended to refer to a variety of art-recognized techniques for introducing foreign nucleic acid (e.g., linear DNA or RNA (e.g., a linearized vector or a gene construct alone without a vector) or nucleic acid in the form of a vector (e.g., a plasmid, phage, phasmid, phagemid,

WO 01/00844 PCT/IB00/00943

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transposon or other DNA) into a host cell, including calcium phosphate or calcium chloride co-precipitation, DEAE-dextran-mediated transfection, lipofection, natural competence, chemical-mediated transfer, or electroporation. Suitable methods for transforming or transfecting host cells can be found in Sambrook, et al. (Molecular Cloning: A Laboratory Manual. 2nd, ed., Cold Spring Harbor Laboratory, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, 1989), and other laboratory manuals.

For stable transfection of mammalian cells, it is known that, depending upon the expression vector and transfection technique used, only a small fraction of cells may integrate the foreign DNA into their genome. In order to identify and select these integrants, a gene that encodes a selectable marker (e.g., resistance to antibiotics) is generally introduced into the host cells along with the gene of interest. Preferred selectable markers include those which confer resistance to drugs, such as G418, hygromycin and methotrexate. Nucleic acid encoding a selectable marker can be introduced into a host cell on the same vector as that encoding an SMP protein or can be introduced on a separate vector. Cells stably transfected with the introduced nucleic acid can be identified by, for example, drug selection (e.g., cells that have incorporated the selectable marker gene will survive, while the other cells die).

To create a homologous recombinant microorganism, a vector is prepared which contains at least a portion of an SMP gene into which a deletion, addition or substitution has been introduced to thereby alter, e.g., functionally disrupt, the SMP gene.

Preferably, this SMP gene is a Corynebacterium glutamicum SMP gene, but it can be a homologue from a related bacterium or even from a mammalian, yeast, or insect source. In a preferred embodiment, the vector is designed such that, upon homologous recombination, the endogenous SMP gene is functionally disrupted (i.e., no longer encodes a functional protein; also referred to as a "knock out" vector). Alternatively, the vector can be designed such that, upon homologous recombination, the endogenous SMP gene is mutated or otherwise altered but still encodes functional protein (e.g., the upstream regulatory region can be altered to thereby alter the expression of the endogenous SMP protein). In the homologous recombination vector, the altered portion of the SMP gene is flanked at its 5' and 3' ends by additional nucleic acid of the SMP gene to allow for homologous recombination to occur between the exogenous SMP gene carried by the vector and an endogenous SMP gene in a microorganism. The additional

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flanking SMP nucleic acid is of sufficient length for successful homologous recombination with the endogenous gene. Typically, several kilobases of flanking DNA (both at the 5' and 3' ends) are included in the vector (see e.g., Thomas, K.R., and Capecchi, M.R. (1987) Cell 51: 503 for a description of homologous recombination vectors). The vector is introduced into a microorganism (e.g., by electroporation) and cells in which the introduced SMP gene has homologously recombined with the endogenous SMP gene are selected, using art-known techniques.

In another embodiment, recombinant microorganisms can be produced which contain selected systems which allow for regulated expression of the introduced gene. For example, inclusion of an SMP gene on a vector placing it under control of the lac operon permits expression of the SMP gene only in the presence of IPTG. Such regulatory systems are well known in the art.

In another embodiment, an endogenous SMP gene in a host cell is disrupted (e.g., by homologous recombination or other genetic means known in the art) such that expression of its protein product does not occur. In another embodiment, an endogenous or introduced SMP gene in a host cell has been altered by one or more point mutations, deletions, or inversions, but still encodes a functional SMP protein. In still another embodiment, one or more of the regulatory regions (e.g., a promoter, repressor, or inducer) of an SMP gene in a microorganism has been altered (e.g., by deletion, truncation, inversion, or point mutation) such that the expression of the SMP gene is modulated. One of ordinary skill in the art will appreciate that host cells containing more than one of the described SMP gene and protein modifications may be readily produced using the methods of the invention, and are meant to be included in the present invention.

A host cell of the invention, such as a prokaryotic or eukaryotic host cell in culture, can be used to produce (i.e., express) an SMP protein. Accordingly, the invention further provides methods for producing SMP proteins using the host cells of the invention. In one embodiment, the method comprises culturing the host cell of invention (into which a recombinant expression vector encoding an SMP protein has been introduced, or into which genome has been introduced a gene encoding a wild-type or altered SMP protein) in a suitable medium until SMP protein is produced. In another

WO 01/00844 PCT/IB00/00943

- 48 -

embodiment, the method further comprises isolating SMP proteins from the medium or the host cell.

C. Isolated SMP Proteins

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Another aspect of the invention pertains to isolated SMP proteins, and biologically active portions thereof. An "isolated" or "purified" protein or biologically active portion thereof is substantially free of cellular material when produced by recombinant DNA techniques, or chemical precursors or other chemicals when chemically synthesized. The language "substantially free of cellular material" includes preparations of SMP protein in which the protein is separated from cellular components of the cells in which it is naturally or recombinantly produced. In one embodiment, the language "substantially free of cellular material" includes preparations of SMP protein having less than about 30% (by dry weight) of non-SMP protein (also referred to herein as a "contaminating protein"), more preferably less than about 20% of non-SMP protein, still more preferably less than about 10% of non-SMP protein, and most preferably less than about 5% non-SMP protein. When the SMP protein or biologically active portion thereof is recombinantly produced, it is also preferably substantially free of culture medium, i.e., culture medium represents less than about 20%, more preferably less than about 10%, and most preferably less than about 5% of the volume of the protein preparation. The language "substantially free of chemical precursors or other chemicals" includes preparations of SMP protein in which the protein is separated from chemical precursors or other chemicals which are involved in the synthesis of the protein. In one embodiment, the language "substantially free of chemical precursors or other chemicals" includes preparations of SMP protein having less than about 30% (by dry weight) of chemical precursors or non-SMP chemicals, more preferably less than about 20% chemical precursors or non-SMP chemicals, still more preferably less than about 10% chemical precursors or non-SMP chemicals, and most preferably less than about 5% chemical precursors or non-SMP chemicals. In preferred embodiments, isolated proteins or biologically active portions thereof lack contaminating proteins from the same organism from which the SMP protein is derived. Typically, such proteins are produced by recombinant expression of, for example, a C. glutamicum SMP protein in a microorganism such as C. glutamicum.

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An isolated SMP protein or a portion thereof of the invention can participate in the metabolism of carbon compounds such as sugars, or in the production of energy compounds (e.g., by oxidative phosphorylation) utilized to drive unfavorable metabolic pathways, or has one or more of the activities set forth in Table 1. In preferred embodiments, the protein or portion thereof comprises an amino acid sequence which is 5 sufficiently homologous to an amino acid sequence of the invention (e.g., a sequence of an even-numbered SEQ ID NO: of the Sequence Listing) such that the protein or portion thereof maintains the ability to perform a function involved in the metabolism of carbon compounds such as sugars or in the generation of energy molecules by processes such as oxidative phosphorylation in Corynebacterium glutamicum. The portion of the protein is preferably a biologically active portion as described herein. In another preferred embodiment, an SMP protein of the invention has an amino acid sequence set forth as an even-numbered SEQ ID NO: of the Sequence Listing. In yet another preferred embodiment, the SMP protein has an amino acid sequence which is encoded by a nucleotide sequence which hybridizes, e.g., hybridizes under stringent conditions, to a nucleotide sequence of the invention (e.g., a sequence of an odd-numbered SEQ ID NO: of the Sequence Listing). In still another preferred embodiment, the SMP protein has an amino acid sequence which is encoded by a nucleotide sequence that is at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, or 60%, preferably at least about 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, or 70%, more preferably at least about 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, or 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, or 90%, or 91%, 92%, 93%, 94%, and even more preferably at least about 95%, 96%, 97%, 98%, 99% or more homologous to one of the nucleic acid sequences of the invention, or a portion thereof. Ranges and identity values intermediate to the above-recited values, (e.g., 70-90% identical or 80-95% identical) are also intended to be encompassed by the present invention. For example, ranges of identity values using a combination of any of the above values recited as upper and/or lower limits are intended to be included. The preferred SMP proteins of the present invention also preferably possess at least one of the SMP activities described herein. For example, a preferred SMP protein of the present invention includes an amino acid sequence encoded by a nucleotide sequence which hybridizes, e.g., hybridizes under stringent conditions, to a nucleotide sequence of the invention, and

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which can perform a function involved in the metabolism of carbon compounds such as sugars or in the generation of energy molecules (e.g., ATP) by processes such as oxidative phosphorylation in *Corynebacterium glutamicum*, or which has one or more of the activities set forth in Table 1.

In other embodiments, the SMP protein is substantially homologous to an amino acid sequence of of the invention (e.g., a sequence of an even-numbered SEQ ID NO: of the Sequence Listing) and retains the functional activity of the protein of one of the amino acid sequences of the invention yet differs in amino acid sequence due to natural variation or mutagenesis, as described in detail in subsection I above. Accordingly, in another embodiment, the SMP protein is a protein which comprises an amino acid sequence which is at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, or 60%, preferably at least about 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, or 70%, more preferably at least about 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, or 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, or 90%, or 91%, 92%, 93%, 94%, and even more preferably at least about 95%, 96%, 97%, 98%, 99% or more homologous to an entire amino acid sequence of the invention and which has at least one of the SMP activities described herein. Ranges and identity values intermediate to the above-recited values, (e.g., 70-90% identical or 80-95% identical) are also intended to be encompassed by the present invention. For example, ranges of identity values using a combination of any of the above values recited as upper and/or lower limits are intended to be included. In another embodiment, the invention pertains to a full length C. glutamicum protein which is substantially homologous to an entire amino acid sequence of the invention.

Biologically active portions of an SMP protein include peptides comprising

amino acid sequences derived from the amino acid sequence of an SMP protein, e.g., an
amino acid sequence of an even-numbered SEQ ID NO: of the Sequence Listing or the
amino acid sequence of a protein homologous to an SMP protein, which include fewer
amino acids than a full length SMP protein or the full length protein which is
homologous to an SMP protein, and exhibit at least one activity of an SMP protein.

Typically, biologically active portions (peptides, e.g., peptides which are, for example,
5, 10, 15, 20, 30, 35, 36, 37, 38, 39, 40, 50, 100 or more amino acids in length) comprise
a domain or motif with at least one activity of an SMP protein. Moreover, other

WO 01/00844 PCT/IB00/00943

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- 51 -

biologically active portions, in which other regions of the protein are deleted, can be prepared by recombinant techniques and evaluated for one or more of the activities described herein. Preferably, the biologically active portions of an SMP protein include one or more selected domains/motifs or portions thereof having biological activity.

SMP proteins are preferably produced by recombinant DNA techniques. For example, a nucleic acid molecule encoding the protein is cloned into an expression vector (as described above), the expression vector is introduced into a host cell (as described above) and the SMP protein is expressed in the host cell. The SMP protein can then be isolated from the cells by an appropriate purification scheme using standard protein purification techniques. Alternative to recombinant expression, an SMP protein, polypeptide, or peptide can be synthesized chemically using standard peptide synthesis techniques. Moreover, native SMP protein can be isolated from cells (e.g., endothelial cells), for example using an anti-SMP antibody, which can be produced by standard techniques utilizing an SMP protein or fragment thereof of this invention.

The invention also provides SMP chimeric or fusion proteins. As used herein, an SMP "chimeric protein" or "fusion protein" comprises an SMP polypeptide operatively linked to a non-SMP polypeptide. An "SMP polypeptide" refers to a polypeptide having an amino acid sequence corresponding to an SMP protein, whereas a "non-SMP polypeptide" refers to a polypeptide having an amino acid sequence corresponding to a protein which is not substantially homologous to the SMP protein, e.g., a protein which is different from the SMP protein and which is derived from the same or a different organism. Within the fusion protein, the term "operatively linked" is intended to indicate that the SMP polypeptide and the non-SMP polypeptide are fused in-frame to each other. The non-SMP polypeptide can be fused to the N-terminus or C-terminus of the SMP polypeptide. For example, in one embodiment the fusion protein is a GST-SMP fusion protein in which the SMP sequences are fused to the C-terminus of the GST sequences. Such fusion proteins can facilitate the purification of recombinant SMP proteins. In another embodiment, the fusion protein is an SMP protein containing a heterologous signal sequence at its N-terminus. In certain host cells (e.g., mammalian host cells), expression and/or secretion of an SMP protein can be increased through use of a heterologous signal sequence.

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Preferably, an SMP chimeric or fusion protein of the invention is produced by standard recombinant DNA techniques. For example, DNA fragments coding for the different polypeptide sequences are ligated together in-frame in accordance with conventional techniques, for example by employing blunt-ended or stagger-ended termini for ligation, restriction enzyme digestion to provide for appropriate termini, filling-in of cohesive ends as appropriate, alkaline phosphatase treatment to avoid undesirable joining, and enzymatic ligation. In another embodiment, the fusion gene can be synthesized by conventional techniques including automated DNA synthesizers. Alternatively, PCR amplification of gene fragments can be carried out using anchor primers which give rise to complementary overhangs between two consecutive gene fragments which can subsequently be annealed and reamplified to generate a chimeric gene sequence (see, for example, Current Protocols in Molecular Biology, Ausubel et al., eds. John Wiley & Sons: 1992). Moreover, many expression vectors are commercially available that already encode a fusion moiety (e.g., a GST polypeptide). An SMP-encoding nucleic acid can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the SMP protein.

Homologues of the SMP protein can be generated by mutagenesis, e.g., discrete point mutation or truncation of the SMP protein. As used herein, the term "homologue" refers to a variant form of the SMP protein which acts as an agonist or antagonist of the activity of the SMP protein. An agonist of the SMP protein can retain substantially the same, or a subset, of the biological activities of the SMP protein. An antagonist of the SMP protein can inhibit one or more of the activities of the naturally occurring form of the SMP protein, by, for example, competitively binding to a downstream or upstream member of the sugar molecule metabolic cascade or the energy-producing pathway which includes the SMP protein.

In an alternative embodiment, homologues of the SMP protein can be identified by screening combinatorial libraries of mutants, e.g., truncation mutants, of the SMP protein for SMP protein agonist or antagonist activity. In one embodiment, a variegated library of SMP variants is generated by combinatorial mutagenesis at the nucleic acid level and is encoded by a variegated gene library. A variegated library of SMP variants can be produced by, for example, enzymatically ligating a mixture of synthetic oligonucleotides into gene sequences such that a degenerate set of potential SMP

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sequences is expressible as individual polypeptides, or alternatively, as a set of larger fusion proteins (e.g., for phage display) containing the set of SMP sequences therein. There are a variety of methods which can be used to produce libraries of potential SMP homologues from a degenerate oligonucleotide sequence. Chemical synthesis of a degenerate gene sequence can be performed in an automatic DNA synthesizer, and the synthetic gene then ligated into an appropriate expression vector. Use of a degenerate set of genes allows for the provision, in one mixture, of all of the sequences encoding the desired set of potential SMP sequences. Methods for synthesizing degenerate oligonucleotides are known in the art (see, e.g., Narang, S.A. (1983) Tetrahedron 39:3; Itakura et al. (1984) Annu. Rev. Biochem. 53:323; Itakura et al. (1984) Science 198:1056; Ike et al. (1983) Nucleic Acid Res. 11:477.

In addition, libraries of fragments of the SMP protein coding can be used to generate a variegated population of SMP fragments for screening and subsequent selection of homologues of an SMP protein. In one embodiment, a library of coding sequence fragments can be generated by treating a double stranded PCR fragment of an SMP coding sequence with a nuclease under conditions wherein nicking occurs only about once per molecule, denaturing the double stranded DNA, renaturing the DNA to form double stranded DNA which can include sense/antisense pairs from different nicked products, removing single stranded portions from reformed duplexes by treatment with S1 nuclease, and ligating the resulting fragment library into an expression vector. By this method, an expression library can be derived which encodes N-terminal, C-terminal and internal fragments of various sizes of the SMP protein.

Several techniques are known in the art for screening gene products of combinatorial libraries made by point mutations or truncation, and for screening cDNA libraries for gene products having a selected property. Such techniques are adaptable for rapid screening of the gene libraries generated by the combinatorial mutagenesis of SMP homologues. The most widely used techniques, which are amenable to high through-put analysis, for screening large gene libraries typically include cloning the gene library into replicable expression vectors, transforming appropriate cells with the resulting library of vectors, and expressing the combinatorial genes under conditions in which detection of a desired activity facilitates isolation of the vector encoding the gene whose product was detected. Recursive ensemble mutagenesis (REM), a new technique which enhances the

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frequency of functional mutants in the libraries, can be used in combination with the screening assays to identify SMP homologues (Arkin and Yourvan (1992) *PNAS* 89:7811-7815; Delgrave *et al.* (1993) *Protein Engineering* 6(3):327-331).

In another embodiment, cell based assays can be exploited to analyze a variegated SMP library, using methods well known in the art.

D. Uses and Methods of the Invention

The nucleic acid molecules, proteins, protein homologues, fusion proteins, primers, vectors, and host cells described herein can be used in one or more of the following methods: identification of *C. glutamicum* and related organisms; mapping of genomes of organisms related to *C. glutamicum*; identification and localization of *C. glutamicum* sequences of interest; evolutionary studies; determination of SMP protein regions required for function; modulation of an SMP protein activity; modulation of the metabolism of one or more sugars; modulation of high-energy molecule production in a cell (*i.e.*, ATP, NADPH); and modulation of cellular production of a desired compound, such as a fine chemical.

The SMP nucleic acid molecules of the invention have a variety of uses. First, they may be used to identify an organism as being Corynebacterium glutamicum or a close relative thereof. Also, they may be used to identify the presence of C. glutamicum or a relative thereof in a mixed population of microorganisms. The invention provides the nucleic acid sequences of a number of C. glutamicum genes; by probing the extracted genomic DNA of a culture of a unique or mixed population of microorganisms under stringent conditions with a probe spanning a region of a C. glutamicum gene which is unique to this organism, one can ascertain whether this organism is present. Although Corynebacterium glutamicum itself is nonpathogenic, it is related to pathogenic species, such as Corynebacterium diphtheriae. Corynebacterium diphtheriae is the causative agent of diphtheria, a rapidly developing, acute, febrile infection which involves both local and systemic pathology. In this disease, a local lesion develops in the upper respiratory tract and involves necrotic injury to epithelial cells; the bacilli secrete toxin which is disseminated through this lesion to distal susceptible tissues of the body. Degenerative changes brought about by the inhibition of protein synthesis in

these tissues, which include heart, muscle, peripheral nerves, adrenals, kidneys, liver and

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spleen, result in the systemic pathology of the disease. Diphtheria continues to have high incidence in many parts of the world, including Africa, Asia, Eastern Europe and the independent states of the former Soviet Union. An ongoing epidemic of diphtheria in the latter two regions has resulted in at least 5,000 deaths since 1990.

In one embodiment, the invention provides a method of identifying the presence or activity of Cornyebacterium diphtheriae in a subject. This method includes detection of one or more of the nucleic acid or amino acid sequences of the invention (e.g., the sequences set forth as odd-numbered or even-numbered SEQ ID NOs, respectively, in the Sequence Listing) in a subject, thereby detecting the presence or activity of Corynebacterium diphtheriae in the subject. C. glutamicum and C. diphtheriae are related bacteria, and many of the nucleic acid and protein molecules in C. glutamicum are homologous to C. diphtheriae nucleic acid and protein molecules, and can therefore be used to detect C. diphtheriae in a subject.

The nucleic acid and protein molecules of the invention may also serve as markers for specific regions of the genome. This has utility not only in the mapping of the genome, but also for functional studies of *C. glutamicum* proteins. For example, to identify the region of the genome to which a particular *C. glutamicum* DNA-binding protein binds, the *C. glutamicum* genome could be digested, and the fragments incubated with the DNA-binding protein. Those which bind the protein may be additionally probed with the nucleic acid molecules of the invention, preferably with readily detectable labels; binding of such a nucleic acid molecule to the genome fragment enables the localization of the fragment to the genome map of *C. glutamicum*, and, when performed multiple times with different enzymes, facilitates a rapid determination of the nucleic acid sequence to which the protein binds. Further, the nucleic acid molecules of the invention may be sufficiently homologous to the sequences of related species such that these nucleic acid molecules may serve as markers for the construction of a genomic map in related bacteria, such as *Brevibacterium lactofermentum*.

The SMP nucleic acid molecules of the invention are also useful for evolutionary and protein structural studies. The metabolic and energy-releasing processes in which the molecules of the invention participate are utilized by a wide variety of prokaryotic and eukaryotic cells; by comparing the sequences of the nucleic acid molecules of the present invention to those encoding similar enzymes from other organisms, the

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evolutionary relatedness of the organisms can be assessed. Similarly, such a comparison permits an assessment of which regions of the sequence are conserved and which are not, which may aid in determining those regions of the protein which are essential for the functioning of the enzyme. This type of determination is of value for protein engineering studies and may give an indication of what the protein can tolerate in terms of mutagenesis without losing function.

Manipulation of the SMP nucleic acid molecules of the invention may result in the production of SMP proteins having functional differences from the wild-type SMP proteins. These proteins may be improved in efficiency or activity, may be present in greater numbers in the cell than is usual, or may be decreased in efficiency or activity.

The invention provides methods for screening molecules which modulate the activity of an SMP protein, either by interacting with the protein itself or a substrate or binding partner of the SMP protein, or by modulating the transcription or translation of an SMP nucleic acid molecule of the invention. In such methods, a microorganism expressing one or more SMP proteins of the invention is contacted with one or more test compounds, and the effect of each test compound on the activity or level of expression of the SMP protein is assessed.

There are a number of mechanisms by which the alteration of an SMP protein of the invention may directly affect the yield, production, and/or efficiency of production of a fine chemical from a *C. glutamicum* strain incorporating such an altered protein. The degradation of high-energy carbon molecules such as sugars, and the conversion of compounds such as NADH and FADH₂ to more useful forms via oxidative phosphorylation results in a number of compounds which themselves may be desirable fine chemicals, such as pyruvate, ATP, NADH, and a number of intermediate sugar compounds. Further, the energy molecules (such as ATP) and the reducing equivalents (such as NADH or NADPH) produced by these metabolic pathways are utilized in the cell to drive reactions which would otherwise be energetically unfavorable. Such unfavorable reactions include many biosynthetic pathways for fine chemicals. By improving the ability of the cell to utilize a particular sugar (e.g., by manipulating the genes encoding enzymes involved in the degradation and conversion of that sugar into energy for the cell), one may increase the amount of energy available to permit

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unfavorable, yet desired metabolic reactions (e.g., the biosynthesis of a desired fine chemical) to occur.

Further, modulation of one or more pathways involved in sugar utilization permits optimization of the conversion of the energy contained within the sugar molecule to the production of one or more desired fine chemicals. For example, by reducing the activity of enzymes involved in, for example, gluconeogenesis, more ATP is available to drive desired biochemical reactions (such as fine chemical biosyntheses) in the cell. Also, the overall production of energy molecules from sugars may be modulated to ensure that the cell maximizes its energy production from each sugar molecule. Inefficient sugar utilization can lead to excess CO₂ production and excess energy, which may result in futile metabolic cycles. By improving the metabolism of sugar molecules, the cell should be able to function more efficiently, with a need for fewer carbon molecules. This should result in an improved fine chemical product: sugar molecule ratio (improved carbon yield), and permits a decrease in the amount of sugars that must be added to the medium in large-scale fermentor culture of such engineered *C. glutamicum*.

The mutagenesis of one or more SMP genes of the invention may also result in SMP proteins having altered activities which indirectly impact the production of one or more desired fine chemicals from C. glutamicum. For example, by increasing the efficiency of utilization of one or more sugars (such that the conversion of the sugar to useful energy molecules is improved), or by increasing the efficiency of conversion of reducing equivalents to useful energy molecules (e.g., by improving the efficiency of oxidative phosphorylation, or the activity of the ATP synthase), one can increase the amount of these high-energy compounds available to the cell to drive normally unfavorable metabolic processes. These processes include the construction of cell walls, transcription, translation, and the biosynthesis of compounds necessary for growth and division of the cells (e.g., nucleotides, amino acids, vitamins, lipids, etc.) (Lengeler et al. (1999) Biology of Prokaryotes, Thieme Verlag: Stuttgart, p. 88-109; 913-918; 875-899). By improving the growth and multiplication of these engineered cells, it is possible to increase both the viability of the cells in large-scale culture, and also to improve their rate of division, such that a relatively larger number of cells can survive in fermentor culture. The yield, production, or efficiency of production may be increased, at least

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due to the presence of a greater number of viable cells, each producing the desired fine chemical.

Further, many of the degradation products produced during sugar metabolism are themselves utilized by the cell as precursors or intermediates for the production of a number of other useful compounds, some of which are fine chemicals. For example, pyruvate is converted into the amino acid alanine, and ribose-5-phosphate is an integral part of, for example, nucleotide molecules. The amount and efficiency of sugar metabolism, then, has a profound effect on the availability of these degradation products in the cell. By increasing the ability of the cell to process sugars, either in terms of efficiency of existing pathways (e.g., by engineering enzymes involved in these pathways such that they are optimized in activity), or by increasing the availability of the enzymes involved in such pathways (e.g., by increasing the number of these enzymes present in the cell), it is possible to also increase the availability of these degradation products in the cell, which should in turn increase the production of many different other desirable compounds in the cell (e.g., fine chemicals).

The aforementioned mutagenesis strategies for SMP proteins to result in increased yields of a fine chemical from *C. glutamicum* are not meant to be limiting; variations on these strategies will be readily apparent to one of ordinary skill in the art. Using such strategies, and incorporating the mechanisms disclosed herein, the nucleic acid and protein molecules of the invention may be utilized to generate *C. glutamicum* or related strains of bacteria expressing mutated SMP nucleic acid and protein molecules such that the yield, production, and/or efficiency of production of a desired compound is improved. This desired compound may be any product produced by *C. glutamicum*, which includes the final products of biosynthesis pathways and intermediates of naturally-occurring metabolic pathways, as well as molecules which do not naturally occur in the metabolism of *C. glutamicum*, but which are produced by a *C. glutamicum* strain of the invention.

This invention is further illustrated by the following examples which should not be construed as limiting. The contents of all references, patent applications, patents, published patent applications, Tables, and the sequence listing cited throughout this application are hereby incorporated by reference.

TABLE 1: GENES IN THE APPLICATION

Function	6-Phosphoglucolactonase L-ribulose-phosphate 4-epimerase RIBULOSE-PHOSPHATE 3-EPIMERASE (EC 5.1.3.1) RIBOSE 5-PHOSPHATE ISOMERASE (EC 5.3.1.6)	Function	SUCCINATE DEHYDROGENASE FLAVOPROTEIN SUBUNIT (EC	1.3.99.1) SUCCINATE DEHYDROGENASE FLAVOPROTEIN SUBUNIT (EC	1.3.99.1) SUCCINATE-SEMIALDEHYDE DEHYDROGENASE (NADP+) (EC 1.2.1.16) SUCCINATE DEHYDROGENASE IRON-SULFUR PROTEIN (EC 1.3.99.1)	FUMARATE HYDRATASE PRECURSOR (EC 4.2.1.2) MALATE DEHYDROGENASE (EC 1.1.1.37) (EC 1.1.1.82) MALATE DEHYDROGENASE (EC 1.1.1.37)			Function	GLUCOKINASE (EC 2.7.1.2) PHOSPHOGLUCOMUTASE (EC 5.4.2.2) / PHOSPHOMANNOMUTASE	(EC 5.4.2.8) PHOSPHOGLUCOMUTASE (EC 5.4.2.2) / PHOSPHOMANNOMUTASE (EC 5.4.2.8)				
NT Stop	15280 3926 14295 5	NT Stop	18785	1614	14015 865	2760 2447 2827			NT Stop	18754 910	657	400	35	5	513
NT Start	14576 4270 13639 346	NT Start	20803	2690	15484	1354 1407 1844		-	NT Start	17786 2571			1624	1588	-
Contig.	VV0074 GR00452 GR00654 GR00290	Contig.	VV0082	GR00380	VV0083 GR00380	GR00131 GR00131 GR00392			Contig.	GR00639 GR00515	VV0086	GR00784	W0043	GR10002	GR00129
Identification Code	RXS02735 RXA01626 RXA02245 RXA01015	Identification Code	RXN01312	F RXA01312	RXN00231 RXA01311 RXA01635	RXA00517 RXA01350	·		Identification Code	RXA02149 RXA01814	RXN02803	F RXA02803	RXN03076	F RXA02854	RXA00511
Amino Acid SEQ ID NO	2 4 9 8	Amino Acid SEQ ID NO	10	12	4 9 8	222		thway	Amino Acid SEQ ID NO	24 26	28	30	32	*	36
Nucleic Acid	1 6 6 7	TCA: Nucleic Acid	o	17	13 13	19 21		EMB-Pathway	Nucleic Acid SEQ ID NO	23 25	27	59	31	33	35

inued)	Function	PHOSPHOGLUCOMUTASE (EC 5.4.2.2) / PHOSPHOMANNOMUTASE	(EC 5.4.2.8) (EC 5.4.2.8)	GLUCOSE-6-PHOSPHATE ISOMERASE (GPI) (EC 5.3.1.9)	GLUCOSE-6-PHOSPHATE ISOMERASE À (GPI A) (EC 5.3.1.9)	PHOSPHOGLYCERATE MUTASE (EC 5.4.2.1)	4 PUDOPTION OF TOTAL OF THE PROPERTY OF THE PR	1-PHOSPHOFRUCTOKINASE (EC.Z.C.1.36) 1-PHOSPHOFRUCTOKINASE (EC.2.C.1.36)	FRUCTOSE-BISPHOSPHATE ALDOLASE (EC 4.1.2.13)	TRIOSEPHOSPHATE ISOMERASE (EC 5.3.1.1)	GLYCERALDEHYDE 3-PHOSPHATE DEHYDROGENASE (EC 1.2.1.12)	GLYCERALDEHYDE-3-PHOSPHATE DEHYDROGENASE HOMOLOG	GLYCERALDEHYDE 3-PHOSPHATE DEHYDROGENASE (EC 1.2.1.12)	PHOSPHOGLYCEKALE KINASE (EC.2.7.2.3)	PYRUVATE KINASE (EC 2.7.1.40)	PHOSPHOENOLPYRUVATE SYNTHASE (EC 2.7.9.2)	PHOSPHOENOLPYRUVATE SYNTHASE (EC 2.7.9.2)	PYRUVATE DEHYDROGENASE (CYTOCHROME) (EC 1.2.2.2)		<u> </u>	•	T ,	PYRUVATE DEHYDROGENASE ET COMPONENT (EC 1.2.4.1)	PYROVATE DEHYDROGENASE ET COMPONENT (EC. 1.2.4.1) PYRUVATE DEHYDROGENASE ET COMPONENT (EC. 1.2.4.1)	. ~	DIHYDROLIPOAMIDE DEHYDROGENASE (EC 1.8.1.4)	DIHYDROLIPOAMIDE DEHYDROGENASE (EC 1.8.1.4)	PHOSPHOENOLPYRUVATE CARBOXYLASE (EC 4.1.1.31)	PYRUVATE CARBOXYLASE (EC 6.4.1.1)	PYRUVATE CARBOXYLASE	PINOVALE CAMBONILAGE (EC 6.4.1.1) PYRIIVATE CARROXYIASE	PYRIVATE CARBOXYLASE (EC 6.4.1.1)	PYRUVATE CARBOXYLASE (EC 6.4.1.1)	MALIC ENZYME (EC 1.1.1.39)						
Table 1 (continued)	NT Stop	103	4	8144	630	2694	2917	846	5813	4 5	2154	366	27227	4943	6741	24935	7004	122	70945	364	4370	3401	5349	20972	325 923	2221	281	955	7650	1362	5	1110	1495	30172	5315	4523	5346	3437	6401	11316
Table	NT Start	1476	897	6525	-	1549	2201	1451	6511	- 6	1165	1397	26451	6382	5302	23934	25155	1552	72801	2	2949	5299	6440	22708	9 e	1391	က	125	2243	<u>-</u>	1291	88	83	27401	4500	5338	305 6305	1842	7783	12539
	Contig.	VV0091	GR00397	GR00014	GR00578	GR00059	GR00720	GR00082	GR00636	GR00032	GR00538	GR00479	GR00654	VV0064	GR00354	GR00654	GK00654	GR00306	8600/	GR00754	GR00755	GR00179	GR00179	VV0135	GR00167	W0019	GR00852	GR00041	VV0049	GK10022	GR10039	VV0047	GR10001	GR00654	VV0047	GK00668	GROOFS GROOFS	W0047	GR00668	VV0079
	Identification Code	RXN01365	F RXA01365	RXA00098	RXA01989	RXA00340	RXA02492	RXA00381	EXA02122	2X 404240	RXA01882	RXA01702	RXA02258	RXN01225	F RXA01225	RXA02256	EXA0225/	RXA01093	RXN02675	F RXA02675	F RXA02695	RXA00682	RXA00683	KXN00635	F RXADD635	RXN03044	F RXA02852	F RXA00268	KXN03086	F KXAU288/ RXN03043	F RXA02897	RXN03083	F RXA02853	RXA02259	RXN02326	F KXA02326	F RX A02327	RXN02328	F RXA02328	RXN01048
	Amino Acid SEQ ID NO	38	40	42	4	46	48	20	52	.	2 80	09	62	64	99	9 9	2 9	4 4	92	78	80	82	84	20 80 80 80 80 80 80 80 80 80 80 80 80 80 80 80 80 80 80 8	8 6	92	94	96	200	36	100	106	108	110	112	114	2 2	120	122	124
	Nucleic Acid SEQ ID NO	37	39	41	43	45	47	49	51	2 3	57	26	61	63	65	29	9 2	73	75	77	62		8	£ 2	5 &	91	93	95	\6 6		103	105	107	109	=======================================	113	117	119	121	123

(penui	Function		MACIO ENZYME (EC 1.1.1.39)	TACTOR FINE (EV. 1.1.38)	C-CACIATE DELYADOOFNADE (EC 1.1.1.27)	D-LACTATE DEHYDDOGENASE (CT IOCHROME) (EC 1.1.2.4)	1.1 ACTATE DEHYDDOCENAGE (CTTOCHROME) (EC 1.1.2.4)	DI ACTATE DEHYDDOGENAGE (CT.OCHROME) (EC 1.1.2.3)	D-LACTATE DEHYDROGENASE (FC 1.1.1.28)	D-LACTATE DEHYDROGENACE (EC 1.1.1.26)	D-3-PHOSPHOGLYCERATE DEHYDROGENASE /EC 111 651	D-3-PHOSPHOGI YOFRATE DEHYDDOGENACE (CO 1.1.1.99)	D-3-PHOSPHOGE YORRATE DEHYDDOGENAGE (FO 1.1.1.93)	D-3-PHOSPHOGE VCENATE DELICATION OF MANAGE (FC 1.1.1.99)	D-3-PHONDHOG! YORKATE DELIVEDOGENAGE (EC. 1.1.39)	IOLB PROTEIN	IOLB PROTEIN: D.FRI ICTORE 1 A.BISDHOSDHATE - CLYOTRONIT OR	PHOSPHATE + D. C. YORDAL DRUYDE S DUDGULATE	JOLS PROTEIN	IOLS PROTEIN	NAGO PROTEIN	PUTATIVE N.GLYCERAL DEHYDE.2. PHOSPHOTEANSES DAYS	GLPX PROTEIN	D-3-PHOSPHOGLYCERATE DEHYDROGENASE (FC 1 1 1 05)	D-3-PHOSPHOGLYCERATE DEHYDROGENASE (EC. 1.1.133)	PHOSPHOGLYCERATE MUTASF /FC 5 4 2 1)	PYRUVATE CARBOXYLASE (EC 6.4.1.1)	PYRUVATE DEHYDROGENASE ET COMPONENT /EC 1 2 4 1	PYRUVATE DEHYDROGENASE E1 COMPONENT (EC 1.2.4.1)	PHOSPHOFNOI DVRIIVATE CAPBOXYKINASE 10707 (50 1.2.4.1)	LIPOAMIDE DEHYDROGENASE COMPONENT (E3) OF BRANCHED.	CHAIN ALPHA-KETO ACID DEHYDROGENASE COMPLEX (EC. 18.1.4) LIPOAMIDE DEHYDROGENASE COMPONENT (E3.) CE BDANCHED	CHAIN ALPHA-KETO ACID DEHYDROGENASE COMPLEX (EC 1.8.1.4)
Table 1 (continued)	NT Stop	000	5655	2820	38606	2837	5417	11666	216	6209	1734	5536	304	. 9	1116	2240	3207		559	562	8298	2074	2989	5224	686	58385	3428	519	281	12541	2296	3533	
Table	NT Start	~	4693	1879	35763	3	4158	9954	-	4611	2645	6138	7	509	568	3127	2344		287	287	7474	1250	3993	6135	1390	59053	3216	310	က	14370	3477	3703	
	Contig.	GROOZGE	GR00046	GR00755	W0176	GR00048	GR00544	W0105	GR00562	GR00562	GR00047	W0157	GR00315	VV0085	GR00316	W0127	GR00239		VV0354	GR00816	VV0019	GR00422	GR00211	VV0213	GR00690	8600	VV0052	W0377	N 0382	W0098	6000A	6000	
	Identification Code	F RXA01048		RXA02694	RXN00296	F RXA00296	RXA01901	RXN01952	F RXA01952	F RXA01955	RXA00293	RXN01130	F RXA01130	RXN03112	F RXA01133	RXN00871	F RXA00871		RXN02829	F RXA02829	RXN01468	F RXA01468.	RXA00794	RXN02920	F RXA02379	RXN02688	RXN03087	RXN03186				RXS01261	
	Amino Acid SEO ID NO	126	128					•	140							154	156			160			166							180		184	
	Nucleic Acid SEQ ID NO	125	127	129	131	133	135	137	139	141	143	145	147	149	151	153	155	!	157	159	161	163	165	16/	60.	ני	1/3	<u>.</u>	//	1/9	181	183	

Glycerol metabolism

	Function	GLYCEROL KINASE (EC 2.7.1.30) GLYCEROL-3-PHOSPHATE DEHYDROGENASE (NAD(P)+) (EC 1.1.1.94) GLYCEROL-3-PHOSPHATE DEHYDROGENASE (NAD(P)+) (EC 1.1.1.94) AEROBIC GLYCEROL-3-PHOSPHATE DEHYDROGENASE (EC 1.1.99.5) GLYCEROL-3-PHOSPHATE REGULON REPRESSOR GLYCEROL-3-PHOSPHATE REGULON REPRESSOR
	NT Stop	2926 4488 1853 1830 2302 147
	NT Start	1400 5483 939 3515 1526 992
	Contig.	GR00749 VV0143 GR00293 GR00525 GR00359 GR00661
	Identification Code	RXA02640 RXN01025 F RXA01025 RXA01851 RXA01242 RXA02288
	Amino Acid SEQ ID NO	188 190 194 194 196
•	Nucleic Acid SEQ ID NO	185 187 191 193

Table 1 (continued)	NT Start NT Stop Function		Ŭ	PRECURSOR	GLYCEROL-3-PHOSPHATE-BINDING PERIPLASMIC PROTEIN	PRECURSOR	Uncharacterized protein involved in glycerol metabolism (homolog of	Drosophila rhomboid)	7 Glycerophosphoryl diester phosphodiesterase	
e 1 (c	NT St		24086		918		3062		22807	
Tabl	NT Start		24949		1736		3808		22091	
	Contig.		W0122		GR00541		GR00703		VV0122	
	Identification Code		RXN01891		F RXA01891		RXA02414		RXN01580	
	Amino Acid	SEQ ID NO	198		200	•	202		204	
	Nucleic Acid	SEQ ID NO	197		199		201		203	

Acetate metabolism

Function	ACETATE KINASE (EC 2.7.2.1)	ACETATE OPERON REPRESSOR	ALCOHOL DEHYDROGENASE (EC 1.1.1.1)	ALDEHYDE DEHYDROGENASE (EC	ALDEHYDE DEHYDROGENASE (EC 1.2.1.3)	ACETOLACTATE SYNTHASE LARGE SUBUNIT (EC 4.1.3.18)	ACETOLACTATE SYNTHASE LARGE SUBUNIT (EC 4.1.3.18)	ACETOLACTATE SYNTHASE LARGE SUBUNIT (EC 4.1.3.18)	ACETOLACTATE SYNTHASE SMALL SUBUNIT (EC 4.1.3.18)							
NT Stop	1357	7941	3391	1959	2419	2945	10159	437	10055	860	3160	14163	320	8254	935	7722
NT Start	2547	8744	4425	1360	1928	3961	11676	108	10678	က	1598	15614	2230	9372	243	8237
Contig.	GR00418	GR00179	GR00037	GR00438	GR00438	GR00498	GR00726	VV0034	VV0155	VV0033	VV0008	VV0315	VV0127	7,0077	VV0264	7,000
Identification Code	RXA01436	RXA00686	RXA00246	RXA01571	RXA01572	RXA01758	RXA02539	RXN03061	RXN03150	RXN01340	RXN01498	RXN02674	RXN00868	RXN01143	RXN01146	RXN01144
Amino Acid	206	208	210	212	214	216	218	220	222	224	226	228	230	232	234	236
Nucleic Acid SEQ ID NO	205	207	209	211	213	215	217	219	221	223	225	227	229	231	233	235

Butanediol, diacetyl and acetoin formation

Function	(S,S)-butane-2,3-diol dehydrogenase (EC 1.1.1.76) ACETOIN(DIACETYL) REDUCTASE (EC 1.1.1.5) ALCOHOL DEHYDROGENASE (EC 1.1.1.1)
NT Stop	7309 5351 28399
NT Start	8082 6103 27383
Contig.	GR00715 GR00710 VV0112
Identification Code	RXA02474 RXA02453 RXS01758
Amino Acid SEQ ID NO	238 240 242
Nucleic Acid SEQ ID NO	237 239 241

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Function	GLUCOSE-6-PHOSPHATE 1-DEHYDROGENASE (EC 1.1.1.49) TRANSALDOLASE (EC 2.2.1.2) TRANSKETOLASE (EC 2.2.1.1) 6-PHOSPHOGI LICONATE DEHYDROGENASE DECABBOXXI ATMOLYCO	1.1.1.44) 6-PHOSPHOGLUCONATE DEHYDROGENASE, DECARBOXYLATING (EC	1.1.1.44) 6-PHOSPHOGLUCONATE DEHYDROGENASE, DECARBOXYLATING (EC 1.1.1.44)
NT Stop	1771 3420 · 4670 510	1366	4448
NT Start	3312 4499 6769 1232	2817	3012
Contig.	GR00763 GR00763 GR00763 GR00270	VV0106	GR00283
Identification Code	RXA02737 RXA02738 RXA02739 RXA00965	RXN00999	F RXA00999
Amino Acid SEQ ID NO	244 246 248 250	252	254
Nucleic Acid SEQ ID NO	243 245 247 249	251	253

Nucleotide sugar conversion

Function	UDP-GALACTOPYRANOSE MUTASE (EC 54.99.9) UDP-GALACTOPYRANOSE MUTASE (EC 54.99.9)	UDP-GALACTOL OF TRANCSE MUTASE (EC 5.4.99.9) UDP-GLUCOSE 6-DEHYDROGENASE (EC 1.1.1.22) UDP-N-ACETYLENOLPYRUVOYLGLUCOSAMINE REDUCTASE (EC	1.1.1.158) UDP-N-ACETYLGLUCOSAMINE PYROPHOSPHORYLASE (EC 2.7.7.23)	UTPGLUCOSE-1-PHOSPHATE URIDYLYLTRANSFERASE (EC 2.7.7.9) UTPGLUCOSE-1-PHOSPHATE LIBIDYLYLTBANSEEDASE (EC 2.7.7.9)	GDP-MANNOSE 6-DEHYDROGENASE (EC 1.1.1.132)	MANNOSE-1-PHOSPHATE GUANYLTRANSFERASE (EC 2.7.7.13) GLICOSE-1-PHOSPHATE ADENIX VITBANSETTASE (FC 2.7.7.13)	GLUCOSE-1-PHOSPHATE THYMINY VI TRANSFERASE (EC 2.7.7.27)	GLUCOSE-1-PHOSPHATE THYMINY VI TRANSEEDASE (EC 2.1.7.24)	GLUCOSE-1-PHOSPHATE THYMINY VI TRANSEEDASE (EC 27.7.24)	D-RIBITOL-5-PHOSPHATE CYTICN VI TRANSCERASE (EC. 2.7.7.24)	DIDP-GLICOSE A R-DEHVDDATAGE (CO 4 2 4 46)
NT Stop	47582 489 5880	3445 3445	1202	130 888	7191	5020 4527	9627	5227	1281	6493	1154
NT Start	48784 1 5383	2345	2302	98/ 573	8351	3301	8848	4448	427	7260	222
Contig.	VV0098 GR00742 GR00749	GR00737 GR00718	GR00352	GR00616	GR00367	GR00626	VV0048	GR00002	GR00438	GR00753	GR00222
Identification Code	RXN02596 F RXA02596 F RXA02642	RXA02572 RXA02485	RXA01216	RXA02028	RXA01262	RXA02063	RXN00014	F RXA00014	RXA01570	RXA02666	RXA00825
Amino Acid SEQ ID NO	256 258 260	262 264	266 268	270	272	276	278	280	282	284	286
c Acid	255 257 259										

Inositol and ribitol metabolism

Function	MYO-INOSITOL 2-DEHYDROGENASE (EC 1.1.1.18)
NT Stop	3209
NT Start	4219
Contig.	GR00539
Identification Code	RXA01887
Amino Acid SEQ ID NO	288
Nucleic Acid SEQ ID NO	287

				anie	able I (conunued	Danii
	Amino Acid	Identification Code	Contig.	NT Start	NT Stop	Function
SEQ ID NO	SEQ ID NO					
	290	RXN00013	VV0048	7966	8838	MYO-INOSITOL-1(OR 4)-MONOPHOSPHATASE 1 (EC 3.1.3.25)
	292	F RXA00013	GR00002	3566	4438	MYO-INOSITOL-1(OR 4)-MONOPHOSPHATASE 1 (EC 3.1.3.25)
	294	RXA01099	GR00306	6328	5504	INOSITOL MONOPHOSPHATE PHOSPHATASE
	296	RXN01332	VV0273	579	4	MYO-INOSITOL 2-DEHYDROGENASE (EC 1.1.1.18)
	298	F RXA01332	GR00388	552	4	MYO-INOSITOL 2-DEHYDROGENASE (EC 1.1.1.18)
	300	RXA01632	GR00454	2338	3342	MYO-INOSITOL 2-DEHYDROGENASE (EC 1.1.1.18)
	302	RXA01633	GR00454	3380	4462	MYO-INOSITOL 2-DEHYDROGENASE (EC 1.1.1.18)
	304	RXN01406	VV0278	2999	1977	MYO-INOSITOL 2-DEHYDROGENASE (EC 1.1.1.18)
	306	RXN01630	VV0050	48113	47037	MYO-INOSITOL 2-DEHYDROGENASE (EC 1.1.1.18)
	308	RXN00528	VV0079	23406	22318	MYO-INOSITOL-1-PHOSPHATE SYNTHASE (EC 5.5.1.4)
	310	RXN03057	VV0028	7017	7688	MYO-INOSITOL 2-DEHYDROGENASE (EC 1.1.1:18)
	312	F RXA02902	GR10040	10277	10948	GLUCOSE-FRUCTOSE OXIDOREDUCTASE PRECURSOR (EC 1.1.99.28)
	314	RXA00251	GR00038	931	224	RIBITOL 2-DEHYDROGENASE (EC 1.1.1.56)

Utilization of sugars

								RECURSOR		RECURSOR					EC (EC		(EC) (EC				
	.47) 1.47)	•			D-RIBOSE-BINDING PERIPLASMIC PROTEIN PRECURSOR			PERIPLASMIC BETA-GLUCOSIDASE/BETA-XYLOSIDASE PRECURSOR		PERIPLASMIC BETA-GLUCOSIDASE/BETA-XYLOSIDASE PRECURSOR		1.67)			GLUCOSE-FRUCTOSE OXIDOREDUCTASE PRECURSOR (EC		GLUCOSEFRUCTOSE OXIDOREDUCTASE PRECURSOR (EC		GLUCOSEFRUCTOSE OXIDOREDUCTASE PRECURSOR (EC		EC 3.2.1.26)	EC 3.2.1.26)	EC 3.2.1.26)
	GLUCOSE 1-DEHYDROGENASE (EC 1.1.1.47) GLUCOSE 1-DEHYDROGENASE II (EC 1.1.1.47)	7.1.12)	7.1.12)	7.1.12)	PLASMIC PROTI	(4:1	(4.1	COSIDASE/BET		COSIDASE/BET		MANNITOL 2-DEHYDROGENASE (EC 1.1.1.67)		se (EC 1.1.1)	XIDOREDUCTA		XIDOREDUCTA		XIDOREDUCTA		SUCROSE-6-PHOSPHATE HYDROLASE (EC 3.2.1.26)	SUCROSE-6-PHOSPHATE HYDROLASE (EC 3.2.1.26)	SUCROSE-6-PHOSPHATE HYDROLASE (EC 3.2.1.26)
	E 1-DEHYDROGIE 1-DEHYDROGI	GLUCONOKINASE (EC 2.7.1.12)	GLUCONOKINASE (EC 2.7.1.12)	GLUCONOKINASE (EC 2.7.1.12)	E-BINDING PERI	FRUCTOKINASE (EC 2.7.1.4)	FRUCTOKINASE (EC 2.7.1.4)	SMIC BETA-GLU	(EC 3.2.1.21) (EC 3.2.1.37)	SMIC BETA-GLU	(EC 3.2.1.21) (EC 3.2.1.37)	OL 2-DEHYDROG	FRUCTOSE REPRESSOR	Hypothetical Oxidoreductase (EC 1.1.1)	E-FRUCTOSE C		E-FRUCTOSE C	_	E-FRUCTOSE (E-6-PHOSPHAT	E-6-PHOSPHAT	E-6-PHOSPHAT
Function	GLUCOSI	GLUCON	GLUCON	GLUCON	D-RIBOSI	FRUCTO	FRUCTO	PERIPLA	(EC 3.2.1	PERIPLA	(EC 3.2.1	MANNITO	FRUCTO	Hypotheti	GLUCOS	1.1.99.28)	GLUCOS	1.1.99.28)	GLUCOS	1.1.99.28)	SUCROS	SUCROS	SUCROS
NT Stop	13090	11114	492	1499	275	5604	1086	56834		1584		10520	7854	8180	5		7050		301		S	9	349
NT Start	12206 7405	9633	1502	1972	1216	6557	565	58477		-		12028	6880	7035	316		6616		735		1246	725	1842
Contig.	VV0090	VV0079	GR00296	GR00296	GR00032	VV0127	GR00240	6000		GR00214		GR00003	GR00725	90000	GR00053		0000		GR00053		GR00007	GR00615	GR00626
Identification Code	RXN02654 F RXA02654	RXN01049	F RXA01049	F RXA01050	RXA00202	RXN00872	F RXA00872	RXN00799		F RXA00799		RXA00032	RXA02528	RXN00316	F RXA00309		RXN00310		F RXA00310		RXA00041	RXA02026	RXA02061
Amino Acid SEQ ID NO	316	320	322	324	326	328	330	332		334		336	338	340	342		344		346		348	350	352
Nucleic Acid SEQ ID NO	315	319	321	323	325	327	329	331		333		335	337	339	341		343		345		347	349	351

(penui	Function		MANNOSE-6-PHOSPHATE ISOMERASE (EC 5.3.1.8)	MANNOSE-6-PHOSPHATE (SOMERASE (EC 5.3.1.8)	INANINOSE-O-PROSPRATE ISOMERASE (EC. 5.3.1.8) 1.4.4. DHA. DI IIOAN BRANDHING ENZYME (EC. 5.3.1.8)	1.4-ALPHA-GILICAN BRANCHING ENZYME (EC. 2.4.1.10)	GLYCOGEN DEBRANCHING ENZYME (EC. 2.4.1.1.19)	GLYCOGEN DEBRANCHING ENZYME (EC 2.4.1.25) (EC 3.2.1.33)	GLYCOGEN OPERON PROTEIN GLGX (EC 3.2.1)	GLYCOGEN PHOSPHORYLASE (EC 2.4.1.1)	ALTICOGEN FROMFROMFILMOR (EC. 2.4.1.1) A) PHA-AMYI AMF (FC. 3.9.1.1)	GLUCOAMYLASE G1 AND G2 PRECURSOR (FC 3 2 1 3)	GLUCOSE-RESISTANCE AMYLASE REGULATOR	XYLULOSE KINASE (EC 2.7.1.17)	XYLULOSE KINASE (EC 2.7.1.17)	RIBOKINASE (EC 2.7.1.15)	RIBOKINASE (EC 2.7.1.15)	RIBOSE OPERON REPRESSOR	6-PHOSPHO-BETA-GLUCOSIDASE (EC 3.2.1.86)	DEOXYRIBOSE-PHOSPHATE ALDOLASE (EC 4.1.2.4)	1-deoxy-D-kylulose 5-phosphate reductoisomerase (EC 1.1.1)	1-DEOXYXYLULOSE-5-PHOSPHATE SYNTHASE	1-DEOXYXYLULOSE-5-PHOSPHATE SYNTHASE	1-DEOXYXYLULOSE-5-PHOSPHATE SYNTHASE	4-ALPHA-GLUCANOTRANSFERASE (EC 2.4.1.25)	4-ALPHA-GLUCANOTRANSFERASE (EC 2.4.1.25), amylomaltase	N-ACETYLASE (EC 3.5.1.25) N ACETYLOLLICOSAMINE & PHOSPHATE BEACETYLASE (EC 3.5.1.25)	N-ACETYL GLIDORAMINY TRANSFERASE (FC 5.9.1.29)	N-ACETYLGLUCOSAMINYLTRANSFERASE (FC 2.4.1.5)	N-ACETYLGLUCOSAMINYLTRANSFERASE (EC 2.4.1)	GLUCOSAMINE-6-PHOSPHATE ISOMERASE (EC 5.3.1.10)	GLUCOSAMINEFRUCTOSE-6-PHOSPHATE AMINOTRANSFERASE	(ISOMERIZING) (EC 2.6.1.16)	URONATE ISOMERASE (EC 5.3.1.12)	URONATE ISOMERASE, Glucuronate isomerase (EC 5.3.1.12)	OKONATE ISOMERASE (EC. 5.3.1.12)	ONOTATION OF A CONTRACT OF A CONTRACT OF A 191	OSENCIOSIDE CACELLEINANSFENASE (EC. 2.3.1.10) D-RIBITOL-5-PHOSPHATE CYTIDYLYLTRANSFERASE (EC. 2.7.7.40)	D-RIBOSE-BINDING PERIPLASMIC PROTEIN PRECURSOR	D-RIBOSE-BINDING PERIPLASMIC PROTEIN PRECURSOR			
Table 1 (continued)	NT Stop		1776	203	1752	3985	1890	1475	17427	16260	1346	2326	920	16532	12352	4923	49244	1118	4	2641	731	2552	5005	1703	3137	1039	1573	6646	3828	L 802	33805	510	547	1279	15397	,	,	4 4	163	2284	6493	275	4258
Table	NT Start		595	50 A		1793	_	ဗ	16981	14749	6 (Ν (15516	10517	4366	50623	က	747	1739	1768	2193	56/6	1094	1230	7	971	8763	592/	3244	35265	1157	1473	2037	17271	c	275	673	672	1611	7260	1216	2097
	Contig.		VV0124	GR00390	GR00743	GR00743	W0184	GR00539	GR00306	W0143	GR00431	VV0318	GRUUGST	GR00639	GR00422	GR00539	W0127	GR00555	GR00762	GR00778	GR00762	GR00729	GROUSES	GROOMS	W0191	GR00436	GR00480	VV0099	GK00242	GROOO?	W0127	GR00520	GR00529	GR00007	GR00422	00000	VVU336 CP10013	GK10015	GR10014	GROOFF	GR00753	GR00032	GR00709
	Identification Code		EXN01369	F RXA01373	RXA02611	RXA02612	RXN01884	F RXA01884	RXA01111	RXN01550	F RXA01550	FX NUZ 100	F RXA02100	RXA02147	RXA01478	RXA01888	RXN01927	F RXA01927	RXA02729	KXA02797	KXA02730	KXA02551	RXA01325	RXA00196	RXN01562	F RXA01562	F RXA01705	KXN008/9	P KARUU6/9	F RX ADDIA3	RXN01752	F RXA01839	RXA01859	RXA00042	RXA01482	0VN00470	F DY A 02872	RXN03180	F RXA02873	RXA02292	RXA02666	RXA00202	RXA02440
	Amino Acid	SEQ ID NO	354 356	358	360	362	364	366	368	370	372	3/4	378	380	382	384	386	388	390	392	394	200	398 400	402	404	406	408	014	1 1 4 1 4 1 4 1 4 1 4 1 4 1 4 1 4 1 4 1	416	418	420	422	424	426	438	430	432	434	436	438	440	442
	Nucleic Acid	SEQ ID NO	353 355	357	359	361	363	365	367	369	371	375	377	379	381	383	385	387	389	186	293	202	369	401	403	405	407	9 4 4	413	5 4	417	419	421	423	425	767	429	431	433	435	437	439	441

A ciolo: M	7.00			Table	Table 1 (continued)	nued)
SEQ ID NO	SEQ ID NO	Identification Code	Contin	N Ottan	NI Stop	Lunction
443	444	RXN01569	6000	41086	4244	dTDP-4-DEHYDRORHAMNOSE REDUCTASE (EC 1.1.1.33)
445	446	F RXA01569	GR00438	2	427	DTDP-4-DEHYDRORHAMNOSE REDUCTASE (EC 1.1.1.133)
447	448	F RXA02055	GR00624	7122	8042	DTDP-4-DEHYDRORHAMNOSE REDUCTASE (EC 1.1.1.133)
449	450	RXA00825	GR00222	222	1154	DTDP-GLUCOSE 4,6-DEHYDRATASE (EC 4.2.1.46)
451	452	EXA02054	GR00624	6103	7119	DTDP-GLUCOSE 4,6-DEHYDRATASE (EC 4.2.1.46)
453	454	KXN00427	W0112	7004	6219	dTDP-RHAMNOSYL TRANSFERASE RFBF (EC 2)
455	456	F RXA00427	GR00098	1591	2022	DTDP-RHAMNOSYL TRANSFERASE RFBF (EC 2)
457	458	RXA00327	GR00057	10263	9880	PROTEIN ARAJ
459	460	RXA00328	GR00057	11147	10656	PROTEIN ARAJ
461	462	RXA00329	GR00057	12390	11167	PROTEIN ARAJ
463	464	RXN01554	W0135	28686	. 26545	GLUCAN ENDO-1,3-BETA-GLUCOSIDASE A1 PRECURSOR (EC 3.2.1.39)
465	466	RXN03015	VV0063	289	œ	UDP-GLUCOSE 6-DEHYDROGENASE (EC 1.1.1.22)
467	468	RXN03056	VV0028	6258	6935	PUTATIVE HEXULOSE-6-PHOSPHATE ISOMERASE (EC 5)
469	470	RXN03030	6000	57006	56443	PERIPLASMIC BETA-GLUCOSIDASE/BETA-XYLOSIDASE PRECURSOR
						(EC 3.2.1.21) (EC 3.2.1.37)
471	472	RXN00401	VV0025	12427	11489	5-DEHYDRO-4-DEOXYGLUCARATE DEHYDRATASE (EC 4.2.1.41)
473	474	RXN02125	VV0102	23242	22442	ALDOSE REDUCTASE (EC 1.1.1.21)
475	476	RXN00200	VV0181	1679	5116	arabinosyl transferase subunit B (EC 2.4.2)
477	478	RXN01175	71007	39688	38303	PHOSPHO-2-DEHYDRO-3-DEOXYHEPTONATE ALDOLASE (EC 4.1.2.15)
479	480	RXN01376	W0091	5610	4750	PUTATIVE GLYCOSYL TRANSFERASE WBIF
481	482	RXN01631	05000	47021	46143	PUTATIVE HEXULOSE-6-PHOSPHATE ISOMERASE (EC 5)
483	484	RXN01593	W0229	13274	12408	NAGD PROTEIN
485	486	RXN00337	W0197	20369	21418	GAI ACTOKINASE (EC. 2.7.1.6)
487	488	RXS00584	10000	5516	6640	CHONOLOGICATION (LO E.T. 1.3) PHONOPHOLO-DEHYDRO-3-DEOXYHEDTONATE ALDOLASE (EC 4 1 2 15)
780	8 6	DXC03674		2	9	DETA LEVORAMINIDACE A DECLIDEDO (CO 3 2 4 E2)
103	, 490 400	DVS02374				DETA-REVOAMMINDANE A PRECORSON (EC. 3.2. 1.32)
-	764	2130000				41 199 28)
493	494	F RXA01915	GR00549	-	1008	GLUCOSE-FRUCTOSE OXIDOREDUCTASE PRECURSOR (EC
						1.1.99.28)
495	496	RXS03224				CYCLOMALTODEXTRINASE (EC 3.2.1.54)
497	498	F RXA00038	GR00006	1417	260	CYCLOMALTODEXTRINASE (EC 3.2.1.54)
499	200	RXC00233				protein involved in sugar metabolism
501	502	RXC00236				Membrane Lipoprotein involved in sugar metabolism
503	504	RXC00271				Exported Protein involved in ribose metabolism
505	206	RXC00338				protein involved in sugar metabolism
507	208	RXC00362				Membrane Spanning Protein involved in metabolism of dipla
509	510	RXC00412				Amino Acid ABC Transporter ATD-Binding Protein involved in sugar
;) :					metabolism
511	512	RXC00526				ABC Transporter ATP-Binding Protein involved in sugar metabolism
513	514	RXC01004				Membrane Spanning Protein involved in sugar metabolism
515	516	RXC01017				Cytosolic Protein involved in sugar metabolism
517	518	RXC01021				Cytosolic Kinase involved in metabolism of sugars and thiamin
519	520	RXC01212				ABC Transporter ATP-Binding Protein involved in sugar metabolism
521	522	RXC01306				Membrane Spanning Protein involved in sugar metabolism
523	524	RXC01366				Cytosolic Protein involved in sugar metabolism
525	526	RXC01372				Cytosolic Protein involved in sugar metabolism

Table 1 (continued		
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Glyoxylate bypass

Function	ISOCITRATE LYASE (EC 4.1.3.1) ISOCITRATE LYASE (EC 4.1.3.1) MALATE SYNTHASE (EC 4.1.3.2) MALATE SYNTHASE (EC 4.1.3.2) GLYOXYLATE-INDUCED PROTEIN GLYOXYLATE-INDUCED PROTEIN	
NT Stop	18365 1773 22475 1663 3958 2430	
NT Start	19708 478 20259 3798 3209	
Contig.	W0176 GR00699 W0176 GR00700 GR00304 GR00539	
Identification Code	RXN02399 F RXA02399 RXN02404 F RXA02404 RXA01089 RXA01886	
Amino Acid SEQ ID NO	590 592 594 596 600	
Nucleic Acid SEQ ID NO	589 591 593 597 599	

Methylcitrate-pathway

Function	2-methylisocitrate synthase (EC 5.3.3)	2-methylisocitrate synthase (EC 5.3.3)	2-methylisocitrate synthase (EC 5.3.3)	2-methylcitrate synthase (EC 4.1.3.31)	2-methylcitrate synthase (EC 4.1.3.31)	2-methylisocitrate synthase (EC 5.3.3)	2-methylcitrate synthase (EC 4.1.3.31)	methylisocitrate lyase (EC 4.1.3.30)	methylisocitrate lyase (EC 4.1.3.30)	LACTOYLGLUTÁTHIONE LYASE (EC 4.4.1.5)			
NT Stop	1576	4	1576	4	2773	6017	901	5	2	764	1815	1902	6266
NT Start	3087	978	1983	621	3069	4647	7	415	209	1906	901	2120	9290
Contig.	VV0092	GR00090	GR00130	GR00130	GR00131	GR00300	W0141	GR00668	GR00669	GR00671	W0141	GR00671	GR00003
Identification Code	RXN03117	F RXA00406	F RXA00514	RXA00512	RXA00518	RXA01077	RXN03144	F RXA02322	RXA02329	RXA02332	RXN02333	F RXA02333	RXA00030
Amino Acid SEQ ID NO	602	604	909	809	610	612	614	616	618	620	622	624	929
Nucleic Acid SEQ ID NO	009	601	603	605	209	609	611	613	615	617	619	621	623

Methyl-Malonyl-CoA-Mutases

Function	METHYLMALONYL-COA MUTASE ALPHA-SUBUNIT (EC 5.4)	METHYLMALONYL-COA MUTASE ALPHA-SUBUNIT (EC 5.4)	METHYLMALONYL-COA MUTASE BETA-SUBUNIT (EC 5.4.9
NT Stop	12059	ည	2009
NT Start	9849	2002	3856
Contig.	W0167	GR00023	GR00023
Identification Code	RXN00148	F RXA00148	RXA00149
Amino Acid SEQ ID NO	628	630	632
Nucleic Acid SEQ ID NO	625	627	629

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Function	PHOSPHOGLYCOLATE PHOSPHATASE (EC 3.1.3.18) PHOSPHOGLYCOLATE PHOSPHATASE (EC 3.1.3.18) PHOSPHOGLYCOLATE PHOSPHATASE (EC 3.1.3.18) PHOSPHOGLYCOLATE PHOSPHATASE (EC 3.1.3.18)		Function	CYTOCHROME D UBIQUINOL OXIDASE SUBUNIT I (EC 1.10.3)	CYTOCHROME DUBIQUINOLOXIDASE SUBUNIT I (EC 1.10.3)	CYTOCHROME C-TYPE BIOGENESIS PROTEIN CON	CYTOCHROME C-TYPE BIOGENESIS PROTEIN CCDA	CYTOCHROME D'UBIQUINOL OXIDASE SUBUNIT II (EC 1.10.3)	CYTOCHROME CONIDASE POLITIEF LIDE (EC. 1.9.5.1) CYTOCHROME COXIDASE SUBLINIT (EC. 1.9.3.1)	CYTOCHROME C OXIDASE POLYPEPTIDE I (EC 1.9.3.1)	CYTOCHROME C OXIDASE POLYPEPTIDE I (EC 1.9.3.1)	CYTOCHROME C OXIDASE POLYPEPTIDE II (EC 1.9.3.1)	CYTOCHROME C OXIDASE POLYPEPTIDE I (EC 1.9.3.1) BIESKE IDON SIII EIID DEOTEIN	PROBABLE CYTOCHROME C OXIDASE ASSEMBLY FACTOR	CYTOCHROME AA3 CONTROLLING PROTEIN	FERREDOXIN	FERREDOXIN	FERREDOXIN VI	FENNEDOMINEMAD(*) REDOCTABE (EC. 1.18.1.3)	ELECTRON TRANSFER FLAVORIO DE META-SUBJINIT	NADH DEHYDROGENASE I CHAIN I (FC. 1.6.4.3)	NADH DEHYDROGENASE I CHAIN I (FC 16.3.3)	NADH DEHYDROGENASE I CHAIN M (EC 1.6.5.3)	NADH DEHYDROGENASE I CHAIN M (EC 1 6 5.3)	NADH DEHYDROGENASE I CHAIN L (EC 1.6.5.3)	NADH DEHYDROGENASE I CHAIN L (EC 1.6.5.3)	NADH-UBIQUINONE OXIDOREDUCTASE CHAIN 2	NADH-UBIQUINONE OXIDOREDUCTASE 39 KD SUBUNIT PRECURSOR (FC 1.6.5.3) (FC 1.6.99.3)	
NT Stop	27532 6 3264 14643		NT Stop	812	- 1690 812	9 9	435	5 29567	4	601	1334	8415	10063 12248	8542	12497	1519	122	2315	24015	24998	9026	1869	7113	3017	2120	3406	43	46287	
NT Start	26879 344 3956 14236		NT Start	2350	2113	212	773	31222	288	1449	1945	7339	11025	7613	13534	1199	436	2302	24965	25783	11299	121	8642	2253	9	2552	846	44824	
Contig.	VV0197 GR00055 GR00645 VV0124		Contig.	VV0174	GR00494	GR00082	GR00083	VV0084	GR00550	GR00717	GR00717	GR00639	GR00639	GR00763	GR00763	GR00355	GR00532	GR00179	GR00032	GR00032	VV0192	GR00160	VV0192	GR00160	GR00249	GR00247	GR00182	08008	
Identification Code	RXN00317 F RXA00317 RXA02196 RXN02461		Identification Code	RXN01744	F RXA01744	RXA00379	RXA00385	RXN02480	F RXA01919	F RXA02480	F RXA02481	EXA02140	RXA02144	RXA02740	RXA02743	RXA01227	RXA01865	RXA00679	RXA00224	RXA00225	RXN00606	F RXA00606	RXN00595	F RXA00608	RXA00913	RXA00909	RXA00700	KXN00483	
Amino Acid SEQ ID NO	634 636 638 640	nain	Amino Acid SEQ ID NO	642	646	648	650 652	654	656	658	099	799	999	668	670	672	676 676	678 678	680	682	684	989	688	069	692	694	969	9A9	
Nucleic Acid SEO ID NO	631 635 637 639	Redox Chain	Nucleic Acid SEQ ID NO	643 643	645	647	649 651	653	655	657	659 664	663	665		699		675			681							693		

(panu	<u>Function</u>	NADH-UBIQUINONE OXIDOREDUCTASE 39 KD SUBUNIT PRECURSOR	(EC. 1.0.0.9) (EC. 1.0.88.9) NADH-DEPENDENT FMN OXYDOREDUCTASE	QUINONE OXIDOREDUCTASE (EC 1.6.5.5)	QUINONE OXIDOREDUCTASE (EC 1.6.5.5)	NADPH-FLAVIN OXIDOREDUCTASE (EC 1.6.99)	NADPH-FLAVIN OXIDOREDUCTASE (EC 1.6.99)	SUCCINATE DEHYDROGENASE IRON-SULFUR PROTEIN (EC 1.3.99.1)	NADH DEHYDROGENASE I CHAIN M (EC 1.6.5.3)	Hydrogenase subunits	NAUH DEHYUKUGENASE (EC 1.6.99.3)	DEHYDROGENANE	FORMATE DEHYDROGENASE ALPHA CHAIN (EC 1.2.1.2)	TOHO PROJEIN	PUHD PROTEIN	CY OCHROME C BIOGENESIS PROTEIN COSA	essential protein similar to cytochrome c	KESC PROTEIN, essential protein similar to cytochrome c biogenesis protein	putative cytochrome oxidase	FLAVOHÉMOPROTEIN / DIHYDROPTERIDINE REDUCTASE (EC	1.6.99.7)	PLAVOHEMOPKO I EIN	GLUTATHIONE STIRANSFERASE (EC. 2.3.1.18) GLITATHIONE DEDENDENT FORMAL DELIVER DELIVERORIASE (FO.	GLOTATIONE-DEPENDENT FORMALDENTOE DEPTOROGENAGE (EU 1211)	OCRC PROTEIN, menaguinol cytochrome c oxidoreductase	NADH DEHYDROGENASE I CHAIN M (EC 1.6.5.3)	NADH-UBIQUINONE OXIDOREDUCTASE CHAIN 4 (EC 1.6.5.3)	Hypothetical Oxidorductase	Hypothetical Oxidoreductase	Hypothetical Oxidoreductase (EC 1.1.1)		Function	ATO CONTUACE A CHAIN CO 2 6 1 24)	ATP SYNTHASE A CHAIN (EC. 5.0.1.54) ATP SYNTHASE A CHAIN (EC. 3.6.1.34)	ATP SYNTHASE ALPHA CHAIN (EC 3.6.1.34)	ATP SYNTHASE BETA CHAIN (EC 3.6.1.34)	ATP SYNTHASE BETA CHAIN (EC. 3.6.1.34) ATP SYNTHASE BETA CHAIN (EC. 3.6.1.34)	
Table 1 (continued)	NT Stop	20569	547	1636	8620	10788	7160	865	368	1259	1) LQ	271	7516	40, 200,	3091)	n	2847	6229	6	31/6	33/3	<u> </u>	11025	4	33063	2794	849	4010		NT Stop	161	1155	2315	3832	3993	
Table	NT Start	19106	1035	2646	9585	9922	6339	1611	1273.	ر د د د	622	7	2556	1119	1291	L 2081	200	410	1876	5602	,	2019	7677	- 607	10138	405	32683	3552	1784	4633		NT Start	1310	394	675	5280 15	3355 3355	
	Contig.	GR00119	GR00427	GR00046	GR00763	W0101	GR00731	GR00380	85000	GR00248	71000	GR00343	GR00183	VV0005	5K00184	VV0025	500000	GK00004	GR00259	W0101	70000	GR00/31	GR00408	12000	GR00639	VV0058	VV0176	VV0317	W0302	VV0101		Contig.	10,000	GR00345	GR00344	W0175	GR00344	
	Identification Code	F RXA00483	RXA01534	RXA00288	RXA02741	RXN02560	F RXA02560	EXA01311	KXN03014	F RXA00910	KANU1895	F KAN1885	KXA00703	FXN00/05	r KXAUU/US	FXNUU388	900000	r rxxu0300	RXA00945	RXN02556		F KXA02556	RX A COROL	000000	RXA02143	RXN03096	RXN02036	RXN02765	RXN02206	RXN02554		Identification Code	DVNO4204	F RX A01204	RXA01201	EXN01193	F RXA01203	
	Amino Acid SEQ ID NO	200	702	704	706	708	710	712	47,	716	120	027	77.	47/	97.	120	7.20	725	734	736	130	2 7 7	740	7	744	746	748	750	752	754	thase	Amino Acid SEQ ID NO	756	758	760	762	766	
	Nucleic Acid SEQ ID NO	669	701	703	705	707	60,	111	51.5	715		2 - 1	12/	57.	C 7 C	171	527	2	733	735	727	130	741	Ē	743	745	747	749	751	753	ATP-Synthase	Nucleic Acid SEQ ID NO	756	757	759	761	765 765	

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(penul	Function	ATP SYNTHASE C CHAIN (EC 3.6.1.34)	ATP SYNTHASE C CHAIN (EC 3.6.1.34)	ATP SYNTHASE DELLA CHAIN (EC.3.6.1.34) ATP SYNTHASE EPSILON CHAIN (EC.3.6.1.34)	ATP SYNTHASE GAMMA CHAIN (EC 3.6.1.34)	ATP-BINDING PROTEIN		Function	CYTOCHROME P450 116 (EC 1.14) Hypothetical Cytochrome c Biogenesis Protein
able 1 (continued	NT Stop	82	318	1141	3349	3274		NT Stop	28581 2004
lable	NT Start	324	139	2 27	2375	4923		NT Start	29864 1150
	Contig.	VV0121	GR00802	GR00343	GR00344	0600/\		Contig.	VV0005 VV0025
	Identification Code	RXN02821	F KXA02821 BXA01200	RXA01194	RXA01202	RXN02434	oolism	Identification Code	RXN00684 RXN00387
	Amino Acid SEQ ID NO	768	22	774	97.	778	Cytochrome metabolism	Amino Acid SEQ ID NO	780 782
	Nucleic Acid SEQ ID NO	767	771	773	775	111	Cytochro	Nucleic Acid SEQ ID NO	779 781

GenBank TM Accession No. A09073 A45579, A45581, A45583, A45587 A45587 A8003132 AB015023 AB018531 AB018531 AB020624 AB023777 AB023777 AB025424 AB0257144 AB027715 AF005242	Gene Name ppg murC; ftsQ; ftsZ dtsR dtsR dtsR acn rep rep acn rep; aad argC	Gene Function Gene Function Phosphoenol pyruvate carboxylase Phosphoenol pyruvate carboxylase Threonine dehydratase Threonine dehydratase Threonine dehydratase Threonine dehydratase Threonine dehydratase Moeckel, B. et micro-organist 951942-A 5 (2383-38 Wachi, M. et a Biotechnol., 5 Kimura, E. et detergent sens lactofermentum Deglutamate racemase transketolase Glutamine 2-oxoglutarate aminotransferase large and small subunits aconitase Replication protein Replication dehydrogenase N-acetylglutamate-5-semialdehydé	Reference Bachmann, B. et al. "DNA fragment coding for phosphoenolpyruvat corboxylase, recombinant DNA carrying said fragment, strains carrying the recombinant DNA and method for producing L-aminino acids using said strains," Patent: EP 0358940-A 3 03/21/90 Moeckel, B. et al. "Production of L-isoleucine by means of recombinant micro-organisms with deregulated threonine dehydratase," Patent: WO 9519442-A 5 07/20/95 gene from coryneform bacteria," Biochem. Biophys. Res. Commun., 236(2):383-388 (1997) Wachi, M. et al. "A murC gene from Coryneform bacteria," Appl. Microbiol. Biotechnol., 51(2):223-228 (1999) Kimura, E. et al. "Molecular cloning of a novel gene, dtsR, which rescues the detergent sensitivity of a mutant derived from Brevibacterium lactofermentum," Biosci. Biotechnol. Biochem., 60(10):1565-1570 (1996)
AF005635	glnA	Glutamine synthetase	
AF030405 AF030520	hisF argG	cyclase Argininosuccinate synthetase	
AF031518	argF	Ornithine carbamolytransferase	
AF038548	arou	3-denydroquinate denydratase Pyruvate carboxylase	

		Table 2 (continued)	inued)
AF038651	dciAE; apt; rel	Dipeptide-binding protein; adenine phosphoribosyltransferase; GTP pyrophosphokinase	Wehmeier, L. et al. "The role of the Corynebacterium glutamicum rel gene in (p)ppGpp metabolism," Microbiology, 144:1853-1862 (1998)
AF041436	argR	Arginine repressor	
AF045998	impA	Inositol monophosphate phosphatase	
AF048764	argH	Argininosuccinate lyase	
AF049897	argC; argJ; argB;	N-acetylglutamylphosphate reductase;	
	argu; argr; argk; argG; argH	ornithine acetyltransferase; N-	
		transminase; ornithine	
		carbamoyltransferase; arginine repressor;	
		argininosuccinate synthase; argininosuccinate Ivase	
AF050109	inhA	Enoyl-acyl carrier protein reductase	
AF050166	hisG	ATP phosphoribosyltransferase	
AF051846	hisA	Phosphoribosylformimino-5-amino-1-	
		phosphoribosyl-4-imidazolecarboxamide	
A DOEDCES		Boinerase	
Ar052652	metA	Homoserine O-acetyltransferase	Park, S. et al. "Isolation and analysis of metA, a methionine biosynthetic gene
			encoding homoserine acetyltransferase in Corynebacterium glutamicum," Mol. Cells., 8(3):286-294 (1998)
AF053071	aroB	Dehydroquinate synthetase	
AF060558	hisH	Glutamine amidotransferase	
AF086704	hisE	Phosphoribosyl-ATP-	
AF114233	aroA	5-enolpyruvylshikimate 3-phosphate	
AF116184	Caca	synthase	
	pani	L-aspartate-alpha-decarboxylase precursor	Dusch, N. et al. "Expression of the Corynebacterium glutamicum panD gene encoding L-aspartate-alpha-decarboxylase leads to pantothenate overproduction in Escherichia coli," <i>Appl. Environ. Microbiol.</i> , 65(4)1530-1530 (1000)
AF124518	aroD; aroE	3-dehydroquinase; shikimate dehydrogenase	
AF124600	aroC; aroK; aroB; pepQ	Chorismate synthase; shikimate kinase; 3-dehydroquinate synthase; putative	
AF145897	inhA		
AF145898	inhA		

		Table 2 (continued)	nued)
AJ001436	ectP	Transport of ectoine, glycine betaine, proline	Peter, H. et al. "Corynebacterium glutamicum is equipped with four secondary carriers for compatible solutes: Identification, sequencing, and characterization of the proline/ectoine uptake system, ProP, and the ectoine/proline/glycine betaine carrier, EctP," J. Bacteriol., 180(22):6005-6012 (1998)
AJ004934	фар	Tetrahydrodipicolinate succinylase (incomplete')	Wehrmann, A. et al. "Different modes of diaminopimelate synthesis and their role in cell wall integrity: A study with Corynebacterium glutamicum," J. Bacteriol., 180(12):3159-3165 (1998)
AJ007732	ppc; secG; amt; ocd; soxA	Phosphoenolpyruvate-carboxylase; ?; high affinity ammonium uptake protein; putative ornithine-cyclodecarboxylase; sarcosine oxidase	
AJ010319	ftsY, glnB, glnD; srp; amtP	Involved in cell division; PII protein; uridylyltransferase (uridylyl-removing enzmye); signal recognition particle; low affinity ammonium uptake protein	Jakoby, M. et al. "Nitrogen regulation in Corynebacterium glutamicum; Isolation of genes involved in biochemical characterization of corresponding proteins," FEMS Microbiol., 173(2):303-310 (1999)
AJ132968	cat	Chloramphenicol aceteyl transferase	
AJ224946	obu	L-malate: quinone oxidoreductase	Molenaar, D. et al. "Biochemical and genetic characterization of the membrane-associated malate dehydrogenase (acceptor) from Corynebacterium glutamicum," Eur. J. Biochem., 254(2):395-403 (1998)
AJ238250	hpu	NADH dehydrogenase	
AJ238703	porA	Porin	Lichtinger, T. et al. "Biochemical and biophysical characterization of the cell wall porin of Corynebacterium glutamicum: The channel is formed by a low molecular mass polypeptide," Biochemistry, 37(43):15024-15032 (1998)
D17429		Transposable element IS31831	Vertes et al. "Isolation and characterization of IS31831, a transposable element from Corynebacterium glutamicum," Mol. Microbiol., 11(4):739-746 (1994)
D84102	odhA	2-oxoglutarate dehydrogenase	Usuda, Y. et al. "Molecular cloning of the Corynebacterium glutamicum (Brevibacterium lactofermentum AJ12036) odhA gene encoding a novel type of 2-oxoglutarate dehydrogenase," <i>Microbiology</i> , 142:3347-3354 (1996)
E01358	hdh; hk		Katsumata, R. et al. "Production of L-thereonine and L-isoleucine," Patent: JP 1987232392-A 1 10/12/87
E01359		Upstream of the start codon of homoserine kinase gene	Katsumata, R. et al. "Production of L-thereonine and L-isoleucine," Patent: JP 1987232392-A 2 10/12/87
E01375		Tryptophan operon	
E01376	trpL; trpE	Leader peptide; anthranilate synthase	Matsui, K. et al. "Tryptophan operon, peptide and protein coded thereby, utilization of tryptophan operon gene expression and production of tryptophan," Patent: JP 1987244382-A 1 10/24/87

	Table 2 (continued)	inued)
E01377	Promoter and operator regions of tryptophan operon	Matsui, K. et al. "Tryptophan operon, peptide and protein coded thereby, utilization of tryptophan operon gene expression and production of tryptophan," Patent: JP 1987244382-A 1 10/24/87
E03937	Biotin-synthase	Hatakeyama, K. et al. "DNA fragment containing gene capable of coding biotin synthetase and its utilization." Patent: JP 1992278088-A 1 10/02/92
E04040	Diamino pelargonic acid aminotransferase	Kohama, K. et al. "Gene coding diaminopelargonic acid aminotransferase and desthiobiotin synthetase and its utilization," Patent: JP 1992330284-A 1 11/18/92
E04041	Desthiobiotinsynthetase	Kohama, K. et al. "Gene coding diaminopelargonic acid aminotransferase and desthiobiotin synthetase and its utilization," Patent: JP 1992330284-A 1 11/18/92
E04307	Flavum aspartase	Kurusu, Y. et al. "Gene DNA coding aspartase and utilization thereof," Patent: JP 1993030977-A 1 02/09/93
E04376		Katsumata, R. et al. "Gene manifestation controlling DNA," Patent: JP 1993056782-A 3 03/09/93
E04377	Isocitric acid Iyase N-terminal fragment	Katsumata, R. et al. "Gene manifestation controlling DNA," Patent: JP 1993056782-A 3 03/09/93
E04484	Prephenate dehydratase	Sotouchi, N. et al. "Production of L-phenylalanine by fermentation," Patent: JP 1993076352-A 2 03/30/93
E05108	Aspartokinase	Fugono, N. et al. "Gene DNA coding Aspartokinase and its use," Patent: JP 1993184366-A 1 07/27/93
E05112	Dihydro-dipichorinate synthetase	Hatakeyama, K. et al. "Gene DNA coding dihydrodipicolinic acid synthetase and its use," Patent: JP 1993184371-A 1 07/27/93
E05776	Diaminopimelic acid dehydrogenase	Kobayashi, M. et al. "Gene DNA coding Diaminopimelic acid dehydrogenase and its use," Patent: JP 1993284970-A 1 11/02/93
E05779	Threonine synthase	Kohama, K. et al. "Gene DNA coding threonine synthase and its use," Patent: JP 1993284972-A 1 11/02/93
E06110	Prephenate dehydratase	Kikuchi, T. et al. "Production of L-phenylalanine by fermentation method," Patent: JP 1993344881-A 1 12/27/93
E06111	Mutated Prephenate dehydratase	Kikuchi, T. et al. "Production of L-phenylalanine by fermentation method," Patent: JP 1993344881-A 1 12/27/93
E06146	Acetohydroxy acid synthetase	Inui, M. et al. "Gene capable of coding Acetohydroxy acid synthetase and its use," Patent: JP 1993344893-A 1 12/27/93
E06825	Aspartokinase	Sugimoto, M. et al. "Mutant aspartokinase gene," patent: JP 1994062866-A 1 03/08/94
E06826	Mutated aspartokinase alpha subunit	Sugimoto, M. et al. "Mutant aspartokinase gene," patent: JP 1994062866-A 1 03/08/94

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nued)	Sugimoto, M. et al. "Mutant aspartokinase gene," patent: JP 1994062866-A 1 03/08/94	Honno, N. et al. "Gene DNA participating in integration of membraneous protein to membrane," Patent: JP 1994169780-A 1 06/21/94	Sato, Y. et al. "Genetic DNA capable of coding Aspartokinase released from feedback inhibition and its utilization," Patent: JP 1994261766-A 1 09/20/94	Sato, Y. et al. "Genetic DNA capable of coding Aspartokinase released from feedback inhibition and its utilization," Patent: JP 1994261766-A 1 09/20/94	Inui, M. et al. "Gene DNA coding acetohydroxy acid isomeroreductase," Patent: JP 1994277067-A 1 10/04/94	Asai, Y. et al. "Gene DNA coding for translocation machinery of protein," Patent: JP 1994277073-A 1 10/04/94	Hatakeyama, K. et al. "DNA fragment having promoter function in coryneform bacterium," Patent: JP 1995031476-A 1 02/03/95	Hatakeyama, K. et al. "DNA fragment having promoter function in coryneform bacterium," Patent: JP 1995031476-A 1 02/03/95	Kohama, K. et al "DNA fragment having promoter function in coryneform bacterium," Patent: JP 1995031478-A 1 02/03/95	Madori, M. et al. "DNA fragment containing gene coding Dihydrodipicolinate acid reductase and utilization thereof," Patent: JP 1995075578-A 1 03/20/95	Madori, M. et al. "DNA fragment containing gene coding Diaminopimelic acid decarboxylase and utilization thereof," Patent: JP 1995075579-A 1 03/20/95	Hatakeyama, K. et al. "Production of L-trypophan," Patent: JP 1997028391-A 1 02/04/97	Moriya, M. et al. "Amplification of gene using artificial transposon," Patent: JP 1997070291-A 03/18/97	Moriya, M. et al. "Amplification of gene using artificial transposon," Patent: JP 1997070291-A 03/18/97	Moriya, M. et al. "Amplification of gene using artificial transposon," Patent: JP 1997070291-A 03/18/97	Moriya, M. et al. "Amplification of gene using artificial transposon," Patent: JP 1997070291-A 03/18/97	Moriya, M. et al. "Amplification of gene using artificial transposon," Patent: JP 1997070291-A 03/18/97
Table 2 (continued)	Mutated aspartokinase alpha subunit	Å	Aspartokinase	Feedback inhibition-released Aspartokinase	Acetohydroxy-acid isomeroreductase	ш	FT aminotransferase and desthiobiotin synthetase promoter region	Biotin synthetase	Aspartase	Dihydrodipicolinate reductase	Diaminopimelic acid decarboxylase	Serine hydroxymethyltransferase	transposase	Arginyl-tRNA synthetase; diaminopimelic acid decarboxylase	Dihydrodipicolinic acid synthetase	aspartokinase	Dihydrodipicolinic acid reductase
		secY				secE	4		_						:		
	E06827	E07701	E08177	E08178, E08179, E08180, E08181, E08182	E08232	E08234	E08643	E08646	E08649	E08900	E08901	E12594	E12760, E12759, E12758	E12764	E12767	E12770	E12773

		Table 2 (continued)	nued)
E13655		ehyd	Hatakeyama, K. et al. "Glucose-6-phosphate dehydrogenase and DNA capable of coding the same," Patent: JP 1997224661-A 1 09/02/97
F01508	livA	Threonine dehydratase	Moeckel, B. et al. "Functional and structural analysis of the threonine dehydratase of Corynebacterium glutamicum," J. Bacteriol., 174:8065-8072 (1992)
L07603	EC 4.2.1.15	3-deoxy-D-arabinoheptulosonate-7- phosphate synthase	Chen, C. et al. "The cloning and nucleotide sequence of Corynebacterium glutamicum 3-deoxy-D-arabinoheptulosonate-7-phosphate synthase gene," FEMS Microbiol. Lett., 107:223-230 (1993)
L09232	IIVB; IIVN; IIVC	Acetohydroxy acid synthase large subunit; Acetohydroxy acid synthase small subunit; Acetohydroxy acid isomeroreductase	Keilhauer, C. et al. "Isoleucine synthesis in Corynebacterium glutamicum: molecular analysis of the ilvB-ilvN-ilvC operon," J. Bacteriol., 175(17):5595-5603 (1993)
L18874	PtsM	Phosphoenolpyruvate sugar phosphotransferase	Fouet, A et al. "Bacillus subtilis sucrose-specific enzyme II of the phosphotransferase system: expression in Escherichia coli and homology to enzymes II from enteric bacteria," PNAS USA, 84(24):8773-8777 (1987); Lee, J.K. et al. "Nucleotide sequence of the gene encoding the Corynebacterium glutamicum mannose enzyme II and analyses of the deduced protein sequence," FEMS Microbiol. Lett., 119(1-2):137-145 (1994)
L27123	aceB	Malate synthase	Lee, H-S. et al. "Molecular characterization of aceB, a gene encoding malate synthase in Corynebacterium glutamicum," J. Microbiol. Biotechnol., 4(4):256-263 (1994)
L27126		Pyruvate kinase	Jetten, M. S. et al. "Structural and functional analysis of pyruvate kinase from Corynebacterium glutamicum," <i>Appl. Environ. Microbiol.</i> , 60(7):2501-2507 (1994)
L28760	aceA	Isocitrate lyase	
L35906	dtxr	Diphtheria toxin repressor	Oguiza, J.A. et al. "Molecular cloning, DNA sequence analysis, and characterization of the Corynebacterium diphtheriae dtxR from Brevibacterium lactofermentum," J. Bacteriol., 177(2):465-467 (1995)
M13774		Prephenate dehydratase	Follettie, M.T. et al. "Molecular cloning and nucleotide sequence of the Corynebacterium glutamicum pheA gene," J. Bacteriol., 167:695-702 (1986)
M16175	5S rRNA		Park, Y-H. et al. "Phylogenetic analysis of the coryneform bacteria by 56 rRNA sequences," J. Bacteriol., 169:1801-1806 (1987)
M16663	trpE	Anthranilate synthase, 5' end	Sano, K. et al. "Structure and function of the trp operon control regions of Brevibacterium lactofermentum, a glutamic-acid-producing bacterium," <i>Gene</i> , 52:191-200 (1987)
M16664	trpA	Tryptophan synthase, 3'end	Sano, K. et al. "Structure and function of the trp operon control regions of Brevibacterium lactofermentum, a glutamic-acid-producing bacterium," <i>Gene</i> , 52:191-200 (1987)

		Table 2 (continued)	ned)
M25819		Phosphoenolpyruvate carboxylase	O'Regan, M. et al. "Cloning and nucleotide sequence of the Phosphoenolpyruvate carboxylase-coding gene of Corynebacterium glutamicum ATCC13032," Gene, 77(2):237-251 (1989)
M85106		23S rRNA gene insertion sequence	Roller, C. et al. "Gram-positive bacteria with a high DNA G+C content are characterized by a common insertion within their 23S rRNA genes," J. Gen. Microbiol., 138:1167-1175 (1992)
M85107, M85108		23S rRNA gene insertion sequence	Roller, C. et al. "Gram-positive bacteria with a high DNA G+C content are characterized by a common insertion within their 23S rRNA genes," J. Gen. Microbiol., 138:1167-1175 (1992)
M89931	aecD; brnQ; yhbw	Beta C-S Iyase; branched-chain amino acid uptake carrier; hypothetical protein yhbw	Rossol, I. et al. "The Corynebacterium glutamicum aecD gene encodes a C-S lyase with alpha, beta-elimination activity that degrades aminoethylcysteine," <i>J. Bacteriol.</i> , 174(9):2968-2977 (1992); Tauch, A. et al. "Isoleucine uptake in Corynebacterium glutamicum ATCC 13032 is directed by the brnQ gene product," <i>Arch. Microbiol.</i> , 169(4):303-312 (1998)
S59299	đ ₁	Leader gene (promoter)	Herry, D.M. et al. "Cloning of the trp gene cluster from a tryptophan-hyperproducing strain of Corynebacterium glutamicum: identification of a mutation in the trp leader sequence," <i>Appl. Environ. Microbiol.</i> , 59(3):791-799 (1993)
U11545	трО	Anthranilate phosphoribosyltransferase	O'Gara, J.P. and Dunican, L.K. (1994) Complete nucleotide sequence of the Corynebacterium glutamicum ATCC 21850 tpD gene." Thesis, Microbiology Department, University College Galway, Ireland.
U13922	cgllM; cglIR; clgIIR	Putative type II 5-cytosoine methyltransferase; putative type II restriction endonuclease; putative type I or type III restriction endonuclease	Schafer, A. et al. "Cloning and characterization of a DNA region encoding a stress-sensitive restriction system from Corynebacterium glutamicum ATCC 13032 and analysis of its role in intergeneric conjugation with Escherichia coli," J. Bacteriol., 176(23):7309-7319 (1994); Schafer, A. et al. "The Corynebacterium glutamicum cglIM gene encoding a 5-cytosine in an McrBC-deficient Escherichia coli strain," Gene, 203(2):95-101 (1997)
U14965 U31224	ррх		Ankri, S. et al. "Mutations in the Corynebacterium glutamicumproline biosynthetic pathway: A natural bypass of the proA step," J. Bacteriol., 178(15):4412-4419 (1996)
U31225	proC	L-proline: NADP+ 5-oxidoreductase	Ankri, S. et al. "Mutations in the Corynebacterium glutamicumproline biosynthetic pathway: A natural bypass of the proA step," J. Bacteriol., 178(15):4412-4419 (1996)
U31230	obg; proB; unkdh	?;gamma glutamyl kinase;similar to D- isomer specific 2-hydroxyacid dehydrogenases	Ankri, S. et al. "Mutations in the Corynebacterium glutamicumproline biosynthetic pathway: A natural bypass of the proA step," J. Bacteriol., 178(15):4412-4419 (1996)

		Table 2 (continued)	ned)
U31281	bioB	Biotin synthase	Serebriiskii, I.G., "Two new members of the bio B superfamily: Cloning, sequencing and expression of bio B genes of Methylobacillus flagellatum and Corynebacterium glutamicum," Gene, 175:15-22 (1996)
U35023	thtR; accBC	Thiosulfate sulfurtransferase; acyl CoA carboxylase	Jager, W. et al. "A Corynebacterium glutamicum gene encoding a two-domain protein similar to biotin carboxylases and biotin-carboxyl-carrier proteins," <i>Arch. Microbiol.</i> , 166(2);76-82 (1996)
U43535	стг	Multidrug resistance protein	Jager, W. et al. "A Corynebacterium glutamicum gene conferring multidrug resistance in the heterologous host Escherichia coli," J. Bacteriol., 179(7):2449-2451 (1997)
U43536	clpB	Heat shock ATP-binding protein	
U53587	aphA-3	3'5"-aminoglycoside phosphotransferase	
U89648		Corynebacterium glutamicum unidentified sequence involved in histidine biosynthesis, partial sequence	
X04960	trpA; trpB; trpC; trpD; trpE; trpG; trpL	Tryptophan operon	Matsui, K. et al. "Complete nucleotide and deduced amino acid sequences of the Brevibacterium lactofermentum tryptophan operon," <i>Nucleic Acids Res.</i> , 14(24):10113-10114 (1986)
X07563	lys A	DAP decarboxylase (meso-diaminopimelate decarboxylase, EC 4.1.1.20)	Yeh, P. et al. "Nucleic sequence of the lysA gene of Corynebacterium glutamicum and possible mechanisms for modulation of its expression," Mol. Gen. Genet., 212(1):112-119 (1988)
X14234	EC 4.1.1.31	Phosphoenolpyruvate carboxylase	Eikmanns, B.J. et al. "The Phosphoenolpyruvate carboxylase gene of Corynebacterium glutamicum: Molecular cloning, nucleotide sequence, and expression," <i>Mol. Gen. Genet.</i> , 218(2):330-339 (1989); Lepiniec, L. et al. "Sorghum Phosphoenolpyruvate carboxylase gene family: structure, function and molecular evolution," <i>Plant. Mol. Biol.</i> , 21 (3):487-502 (1993)
X17313	fda	Fructose-bisphosphate aldolase	Von der Osten, C.H. et al. "Molecular cloning, nucleotide sequence and fine- structural analysis of the Corynebacterium glutamicum fda gene: structural comparison of C. glutamicum fructose-1, 6-biphosphate aldolase to class I and class II aldolases," Mol. Microbiol.
X53993	dapA	L-2, 3-dihydrodipicolinate synthetase (EC 4.2.1.52)	Bonnassie, S. et al. "Nucleic sequence of the dapA gene from Corynebacterium glutamicum," Nucleic Acids Res., 18(21):6421 (1990)
X54223		AttB-related site	Cianciotto, N. et al. "DNA sequence homology between att B-related sites of Corynebacterium diphtheriae, Corynebacterium ulcerans, Corynebacterium glutamicum, and the attP site of lambdacorynephage," FEMS. Microbiol, Lett., 66:299-302 (1990)
X54740	argS; lysA	Arginyl-tRNA synthetase; Diaminopimelate decarboxylase	Marcel, T. et al. "Nucleotide sequence and organization of the upstream region of the Corynebacterium glutamicum lysA gene," Mol. Microbiol., 4(11):1819-1830 (1990)

		Table 2 (continued)	nued)
X55994	trpL; trpE	ide; anth	Heery, D.M. et al. "Nucleotide sequence of the Corynebacterium glutamicum trpE gene," Nucleic Acids Res., 18(23):7138 (1990)
X56037	thrC	Threonine synthase	Han, K.S. et al. "The molecular structure of the Corynebacterium glutamicum threonine synthase gene," <i>Mol. Microbiol.</i> , 4(10):1693-1702 (1990)
X56075	attB-related site	Attachment site	Cianciotto, N. et al. "DNA sequence homology between att B-related sites of Corynebacterium diphtheriae, Corynebacterium ulcerans, Corynebacterium glutamicum, and the attP site of lambdacorynephage," FEMS. Microbiol, Lett., 66:299-302 (1990)
X57226	lysC-alpha; lysC-beta; asd	Aspartokinase-alpha subunit; Aspartokinase-beta subunit; aspartate beta semialdehyde dehydrogenase	Kalinowski, J. et al. "Genetic and biochemical analysis of the Aspartokinase from Corynebacterium glutamicum," <i>Mol. Microbiol.</i> , 5(5):1197-1204 (1991); Kalinowski, J. et al. "Aspartokinase genes lysC alpha and lysC beta overlap and are adjacent to the aspertate beta-semialdehyde dehydrogenase gene asd in Corynebacterium glutamicum," <i>Mol. Gen. Genet.</i> , 224(3):317-324 (1990)
X59403	gap;pgk; tpi	Glyceraldehyde-3-phosphate; phosphoglycerate kinase; triosephosphate isomerase	Eikmanns, B.J. "Identification, sequence analysis, and expression of a Corynebacterium glutamicum gene cluster encoding the three glycolytic enzymes glyceraldehyde-3-phosphate dehydrogenase, 3-phosphoglycerate kinase, and triosephosphate isomeras," J. Bacteriol., 174(19):6076-6086 (1992)
X59404	dpb	Glutamate dehydrogenase	Bormann, E.R. et al. "Molecular analysis of the Corynebacterium glutamicum gdh gene encoding glutamate dehydrogenase," Mol. Microbiol., 6(3):317-326 (1992)
X60312	lysi	L-lysine permease	Seep-Feldhaus, A.H. et al. "Molecular analysis of the Corynebacterium glutamicum lysl gene involved in lysine uptake," Mol. Microbiol., 5(12):2995-3005 (1991)
X66078	cop1	Ps1 protein	Joliff, G. et al. "Cloning and nucleotide sequence of the csp1 gene encoding PS1, one of the two major secreted proteins of Corynebacterium glutamicum: The deduced N-terminal region of PS1 is similar to the Mycobacterium antigen 85 complex," Mol. Microbiol., 6(16):2349-2362 (1992)
X66112	glt	Citrate synthase	Eikmanns, B.J. et al. "Cloning sequence, expression and transcriptional analysis of the Corynebacterium glutamicum gltA gene encoding citrate synthase," <i>Microbiol.</i> , 140:1817-1828 (1994)
X67737	dapB	Dihydrodipicolinate reductase	
X69103	csp2	Surface layer protein PS2	Peyret, J.L. et al. "Characterization of the cspB gene encoding PS2, an ordered surface-layer protein in Corynebacterium glutamicum," Mol. Microbiol., 9(1):97-109 (1993)
X69104		IS3 related insertion element	Bonamy, C. et al. "Identification of IS1206, a Corynebacterium glutamicum IS3-related insertion sequence and phylogenetic analysis," Mol. Microbiol., 14(3):571-581 (1994)

		Table 2 (continued)	inued)
X70959	leuA	Isopropylmalate synthase	Patek, M. et al. "Leucine synthesis in Corynebacterium glutamicum: enzyme activities, structure of leuA, and effect of leuA inactivation on lysine synthesis." Appl. Environ. Microbiol. 60(1):133-140 (1994)
X71489	icd	Isocitrate dehydrogenase (NADP+)	Eikmanns, B.J. et al. "Cloning sequence analysis, expression, and inactivation of the Corynebacterium glutamicum icd gene encoding isocitrate dehydrogenase and biochemical characterization of the enzyme," J. Bacteriol., 177(3):774-782 (1995)
X72855	GDHA	Glutamate dehydrogenase (NADP+)	
X75083, X70584	mtrA	5-methyltryptophan resistance	Heery, D.M. et al. "A sequence from a tryptophan-hyperproducing strain of Corynebacterium glutamicum encoding resistance to 5-methyltryptophan," Biochem Biochem Biochem Biochem
X75085	recA		Fitzpatrick, R. et al. "Construction and characterization of recA mutant strains of Corynebacterium glutamicum and Brevibacterium lactofermentum," Appl. Microhiol Riotechnol 42(4):575-580 (1994)
X75504	aceA; thiX	Partial Isocitrate lyase; ?	Reinscheid, D.J. et al. "Characterization of the isocitrate lyase gene from Corynebacterium glutamicum and biochemical analysis of the enzyme," J. Bacteriol. 176(12):3474-3483 (1994)
X76875		ATPase beta-subunit	Ludwig, W. et al. "Phylogenetic relationships of bacteria based on comparative sequence analysis of elongation factor Tu and ATP-synthase beta-subunit genes." Antonie Van Leeuwenhoek 64-285-305 (1993)
X77034	tuf	Elongation factor Tu	Ludwig, W. et al. "Phylogenetic relationships of bacteria based on comparative sequence analysis of elongation factor Tu and ATP-synthase beta-subunit genes." Antonie Van Leeuwenhoek, 64-285-305 (1993)
X77384	recA		Billman-Jacobe, H. "Nucleotide sequence of a recA gene from Corynebacterium glutamicum." DNA Sea. 4(6):403-404 (1994)
X78491	aceB	Malate synthase	Reinscheid, D.J. et al. "Malate synthase from Corynebacterium glutamicum pta-ack operon encoding phosphotransacetylase: sequence analysis," <i>Microbiology</i> , 140-3099-3108 (1994)
X80629	I6S rDNA	16S ribosomal RNA	Rainey, F.A. et al. "Phylogenetic analysis of the genera Rhodococcus and Norcardia and evidence for the evolutionary origin of the genus Norcardia from within the radiation of Rhodococcus species," <i>Microbiol.</i> , 141:523-528 (1995)
16118X	gluA; gluB; gluC; gluD	Glutamate uptake system	Kronemeyer, W. et al. "Structure of the gluABCD cluster encoding the glutamate uptake system of Corynebacterium glutamicum," J. Bacteriol., 177(5):1152-1158 (1995)
X81379	dapE	Succinyldiaminopimelate desuccinylase	Wehrmann, A. et al. "Analysis of different DNA fragments of Corynebacterium glutamicum complementing dapE of Escherichia coli," <i>Microbiology</i> , 40:3349-56 (1994)

		Table 2 (continued)	nued)
X82061	16S rDNA	16S ribosomal RNA	Ruimy, R. et al. "Phylogeny of the genus Corynebacterium deduced from analyses of small-subunit ribosomal DNA sequences," Int. J. Syst. Bacteriol., 45(4):740-746 (1995)
X82928	asd; lysC	Aspartate-semialdehyde dehydrogenase; ?	Serebrijski, I. et al. "Multicopy suppression by asd gene and osmotic stress-dependent complementation by heterologous proA in proA mutants," J. Bacteriol., 177(24):7255-7260 (1995)
X82929	proA	Gamma-glutamyl phosphate reductase	Serebrijski, I. et al. "Multicopy suppression by asd gene and osmotic stress-dependent complementation by heterologous proA in proA mutants," J. Bacteriol., 177(24):7255-7260 (1995)
X84257	16S rDNA	16S ribosomal RNA	Pascual, C. et al. "Phylogenetic analysis of the genus Corynebacterium based on 16S rRNA gene sequences," Int. J. Syst. Bacteriol., 45(4):724-728 (1995)
X85965	aroP; dapE	Aromatic amino acid permease; ?	Wehrmann et al. "Functional analysis of sequences adjacent to dapE of C. glutamicum proline reveals the presence of aroP, which encodes the aromatic amino acid transporter," J. Bacteriol., 177(20):5991-5993 (1995)
X86157	argB; argC; argD; argF; argJ	Acetylglutamate kinase; N-acetyl-gamma- glutamyl-phosphate reductase; acetylornithine aminotransferase; ornithine carbamoyltransferase; glutamate N- acetyltransferase	Sakanyan, V. et al. "Genes and enzymes of the acetyl cycle of arginine biosynthesis in Corynebacterium glutamicum: enzyme evolution in the early steps of the arginine pathway," <i>Microbiology</i> , 142:99-108 (1996)
X89084	pta; ackA	Phosphate acetyltransferase; acetate kinase	Reinscheid, D.J. et al. "Cloning, sequence analysis, expression and inactivation of the Corynebacterium glutamicum pta-ack operon encoding phosphotransacetylase and acetate kinase," <i>Microbiology</i> , 145:503-513 (1999)
X89850	attB	Attachment site	Le Marrec, C. et al. "Genetic characterization of site-specific integration functions of phi AAU2 infecting "Arthrobacter aureus C70," J. Bacteriol., 178(7):1996-2004 (1996)
X90356		Promoter fragment F1	Patek, M. et al. "Promoters from Corynebacterium glutamicum: cloning, molecular analysis and search for a consensus motif," Microbiology, 142:1297-1309 (1996)
X90357		Promoter fragment F2	Patek, M. et al. "Promoters from Corynebacterium glutamicum: cloning, molecular analysis and search for a consensus motif," <i>Microbiology</i> , 142:1297-1309 (1996)
X90358		Promoter fragment F10	Patek, M. et al. "Promoters from Corynebacterium glutamicum: cloning, molecular analysis and search for a consensus motif," <i>Microbiology</i> , 142:1297-1309 (1996)
X90359		Promoter fragment F13	Patek, M. et al. "Promoters from Corynebacterium glutamicum: cloning, molecular analysis and search for a consensus motif," <i>Microbiology</i> , 142:1297-1309 (1996)

		Table 2 (continued)	nued)
X90360		Promoter fragment F22	Datak M at al "Dromotor from Commohants
			molecular analysis and search for a consensus motif." Microbioloφ.
17000			142:1297-1309 (1996)
A90361		Promoter fragment F34	Patek, M. et al. "Promoters from Corynebacterium glutamicum: cloning,
			molecular analysis and search for a consensus motif," Microbiology,
X90362		Promoter fragment F37	Patek M et al "Demoteer from California"
			and search for a consensus motif." Microbiology, 142:1297-1309 (1996)
X90363		Promoter fragment F45	Patek, M. et al. "Promoters from Corynebacterium glutamicum: cloning,
			molecular analysis and search for a consensus motif," Microbiology, 142:1297-1309 (1996)
X90364		Promoter fragment F64	Patek, M. et al. "Promoters from Corynebacterium glutamicum: cloning
			molecular analysis and search for a consensus motif," Microbiology, 142:1297-1309 (1996)
X90365		Promoter fragment F75	Patek, M. et al. "Promoters from Corynebacterium glutamicum: cloning.
			molecular analysis and search for a consensus motif," Microbiology,
V00355			142:1297-1309 (1996)
99506V		Promoter fragment PF101	Patek, M. et al. "Promoters from Corynebacterium glutamicum: cloning,
	·		molecular analysis and search for a consensus motif," Microbiology,
X90367		Promoter fragment PF104	Patek M et al "Promoters from Convenencium alutominum: alonino
)	molecular analysis and search for a consensus motif." Microbiology
7007			142:1297-1309 (1996)
896088		Promoter fragment PF109	Patek, M. et al. "Promoters from Corynebacterium glutamicum: cloning,
			molecular analysis and search for a consensus motif," Microbiology, 142:1297-1309 (1996)
X93513	amt	Ammonium transport system	Siewe, R.M. et al. "Functional and genetic characterization of the (methyl)
			ammonium uptake carrier of Corynebacterium glutamicum," J. Biol. Chem., 271(10):5398-5403 (1996)
X93514	betP	Glycine betaine transport system	Peter, H. et al. "Isolation, characterization, and expression of the
			Corynebacterium glutamicum betP gene, encoding the transport system for the
X05640	l'Jac		compatible solute glycine betaine," J. Bacteriol., 178(17):5229-5234 (1996)
2000	<u> </u>		Patek, M. et al. "Identification and transcriptional analysis of the dapB-ORF2-
			wapha-Oki's operou of Corynebacterium glutamicum, encoding two enzymes involved in L-lysine synthesis," <i>Biotechnol. Lett.</i> , 19:1113-1117 (1997)
X96471	lysE; lysG	Lysine exporter protein; Lysine export	Vrljic, M. et al. "A new type of transporter with a new type of cellular
		regulator protein	function: L-lysine export from Corynebacterium glutamicum," Mol.
			Microbiol., 22(5):815-826 (1996)

		Table 2 (continued)	(pənu
X96580	panB; panC; xylB	3-methyl-2-oxobutanoate hydroxymethyltransferase; pantoate-beta-alanine ligase; xylulokinase	Sahm, H. et al. "D-pantothenate synthesis in Corynebacterium glutamicum and use of panBC and genes encoding L-valine synthesis for D-pantothenate overproduction," Appl. Environ. Microbiol., 65(5):1973-1979 (1999)
X96962		Insertion sequence IS1207 and transposase	
X99289		Elongation factor P	Ramos, A. et al. "Cloning, sequencing and expression of the gene encoding elongation factor P in the amino-acid producer Brevibacterium lactofermentum (Corynebacterium glutamicum ATCC 13869)," Gene, 198:217-222 (1997)
Y00140	thrB	Homoserine kinase	Mateos, L.M. et al. "Nucleotide sequence of the homoserine kinase (thrB) gene of the Brevibacterium lactofermentum," Nucleic Acids Res., 15(9):3922 (1987)
Y00151	qpp	Meso-diaminopimelate D-dehydrogenase (EC 1.4.1.16)	Ishino, S. et al. "Nucleotide sequence of the meso-diaminopimelate D-dehydrogenase gene from Corynebacterium glutamicum," Nucleic Acids Res., 15(9):3917 (1987)
Y00476	thrA	Homoserine dehydrogenase	Mateos, L.M. et al. "Nucleotide sequence of the homoserine dehydrogenase (thrA) gene of the Brevibacterium lactofermentum," Nucleic Acids Res., 15(24):10598 (1987)
Y00546	hom; thrB	Homoserine dehydrogenase; homoserine kinase	Peoples, O.P. et al. "Nucleotide sequence and fine structural analysis of the Corynebacterium glutamicum hom-thrB operon," <i>Mol. Microbiol.</i> , 2(1):63-72 (1988)
Y08964	murC; ftsQ/divD; ftsZ	UPD-N-acetylmuramate-alanine ligase; division initiation protein or cell division protein; cell division protein	Honrubia, M.P. et al. "Identification, characterization, and chromosomal organization of the ftsZ gene from Brevibacterium lactofermentum," Mol. Gen. Genet., 259(1):97-104 (1998)
Y09163	putP	High affinity proline transport system	Peter, H. et al. "Isolation of the putP gene of Corynebacterium glutamicumproline and characterization of a low-affinity uptake system for compatible solutes," <i>Arch. Microbiol.</i> , 168(2):143-151 (1997)
Y09548	pyc	Pyruvate carboxylase	Peters-Wendisch, P.G. et al. "Pyruvate carboxylase from Corynebacterium glutamicum: characterization, expression and inactivation of the pyc gene," <i>Microbiology</i> , 144:915-927 (1998)
Y09578	leuB	3-isopropylmalate dehydrogenase	Patek, M. et al. "Analysis of the leuB gene from Corynebacterium glutamicum," Appl. Microbiol. Biotechnol., 50(1):42-47 (1998)
Y12472		Attachment site bacteriophage Phi-16	Moreau, S. et al. "Site-specific integration of corynephage Phi-16: The construction of an integration vector," Microbiol., 145:539-548 (1999)
Y12537	proP	Proline/ectoine uptake system protein	Peter, H. et al. "Corynebacterium glutamicum is equipped with four secondary carriers for compatible solutes: Identification, sequencing, and characterization of the proline/ectoine uptake system, ProP, and the ectoine/proline/glycine
			betaine carrier, EctP," J. Bacieriol., 180(22):6005-6012 (1998)

		Table 2 (continued)	nned)
Y13221	glnA	Glutamine synthetase I	Jakoby, M. et al. "Isolation of Corynebacterium glutamicum glnA gene
Y16642	pdl	Dihydrolipoamide dehydrogenase	COCCUIE ELIGIBILITIES 1, FEMS MICROSIOL LEIL, 154(1):81-88 (1997)
Y18059		Attachment site Corynephage 304L	Moreau, S. et al. "Analysis of the integration functions of φ304L: An integrase module among convenhance." Virology, 255(1),150,150,150,150,150,150,150,150,150,150
Z21501	argS; lysA	Arginyl-tRNA synthetase; diaminopimelate decarboxylase (partial)	Oguiza, J.A. et al. "A gene encoding arginyl-tRNA synthetase is located in the upstream region of the lysA gene in Brevibacterium lactofermentum: Regulation of argS-lysA cluster expression by arginine," J. Barteriol 175(22):7356-710033
Z21502	dapA; dapB	Dihydrodipicolinate synthase; dihydrodipicolinate reductase	Pisabarro, A. et al. "A cluster of three genes (dapA, orf2, and dapB) of Brevibacterium lactofermentum encodes dihydrodipicolinate reductase, and a third polypeptide of unknown function," J. Bacieriol., 175(9):2743-2749
Z29563	thrC	Threonine synthase	Malumbres, M. et al. "Analysis and expression of the thrC gene of the encoded threoning synthese," And Emison Minerally (20/2020)
Z46753	16S rDNA	Gene for 16S ribosomal RNA	
Z 49822	sigA	SigA sigma factor	Oguiza, J.A. et al "Multiple sigma factor genes in Brevibacterium lactofermentum: Characterization of sigA and sigB," J. Bacteriol., 178(2):550-553 (1996)
Z49823	galE; dtxR	Catalytic activity UDP-galactose 4- epimerase; diphtheria toxin regulatory protein	Oguiza, J.A. et al "The galE gene encoding the UDP-galactose 4-epimerase of Brevibacterium lactofermentum is coupled transcriptionally to the dmdR gene." Gene. 177:103-107 (1996)
Z49824	orfl; sigB	?; SigB sigma factor	Oguiza, J.A. et al "Multiple sigma factor genes in Brevibacterium lactofermentum: Characterization of sigA and sigB," J. Bacteriol., 178(2):550-553 (1996)
266534		Transposase	Correia, A. et al. "Cloning and characterization of an IS-like element present in the genome of Brevibacterium lactofermentum ATCC 13869," Gene, 170(1):91-94 (1996)
A sequence for the published ver.	this gene was published in sion. It is believed that the	the indicated reference. However, the sequence published version relied on an incorrect start co	A sequence for this gene was published in the indicated reference. However, the sequence obtained by the inventors of the present application is significantly longer than the published version. It is believed that the published version relied on an incorrect start codon, and thus represents only a fragment of the actual coding region.

TABLE 3: Corynebacterium and Brevibacterium Strains Which May be Used in the Practice of the Invention

Conjust	species : 2	ATCC	FFRM	ENRRIS.	СЕСТ	NEIMB	_CBS	NCTE	DSMZ
The state of the s	the winter the second	21054			7			- Table	
Brevibacterium	ammoniagenes	19350			ļ				
Brevibacterium	ammoniagenes	19351			<u> </u>				
Brevibacterium	ammoniagenes	l	ļ		<u> </u>				
Brevibacterium	ammoniagenes	19352		ļ	ļ				
Brevibacterium	ammoniagenes	19353							
Brevibacterium	ammoniagenes	19354							
Brevibacterium	ammoniagenes	19355							
Brevibacterium	ammoniagenes	19356						<u> </u>	
Brevibacterium	ammoniagenes	21055							
Brevibacterium	ammoniagenes	21077			ļ			<u> </u>	
Brevibacterium	ammoniagenes	21553			<u> </u>				
Brevibacterium	ammoniagenes	21580	<u> </u>						
Brevibacterium	ammoniagenes	39101						<u> </u>	
Brevibacterium	butanicum	21196		<u></u>	<u> </u>			ļ	
Brevibacterium	divaricatum	21792	P928						
Brevibacterium	flavum	21474						L	
Brevibacterium	flavum	21129							
Brevibacterium	flavum	21518							
Brevibacterium	flavum			B11474					
Brevibacterium	flavum			B11472					L
Brevibacterium	flavum	21127							
Brevibacterium	flavum	21128							
Brevibacterium	flavum	21427							
Brevibacterium	flavum	21475							
Brevibacterium	flavum	21517							
Brevibacterium	flavum	21528							
Brevibacterium	flavum	21529							
Brevibacterium	flavum			B11477					
Brevibacterium	flavum		<u> </u>	B11478					
Brevibacterium	flavum	21127							
Brevibacterium	flavum			B11474					
Brevibacterium	healii	15527	1						
Brevibacterium	ketoglutamicum	21004							
Brevibacterium	ketoglutamicum	21089			1	<u> </u>			
Brevibacterium	ketosoreductum	21914	 		 				
Brevibacterium	lactofermentum	 	 		70				
Brevibacterium	lactofermentum		 	1	74				
Brevibacterium	lactofermentum	 	1	<u> </u>	77				1
Brevibacterium	lactofermentum	21798	 		†	1			<u> </u>
Brevibacterium	lactofermentum	21799	 		 	 	i	†	
Brevibacterium	lactofermentum	21800		 	—	 		 	
Brevibacterium	lactofermentum	21801	 	 	+		 	 	
Brevibacterium	lactofermentum	21001	 	B11470	 			 	
	·	 	 	B11471		┼──	 	+	
Brevibacterium	lactofermentum	l	<u> </u>	10114/1	L	L		<u> </u>	<u> </u>

Genus			FERM	NRRL	CECT	NCIMB	CBS	NCTC	DSMZ
Brevibacterium	lactofermentum	21086							
Brevibacterium	lactofermentum	21420			T				
Brevibacterium	lactofermentum	21086			T				
Brevibacterium	lactofermentum	31269		<u> </u>	1			 	
Brevibacterium	linens	9174							
Brevibacterium	linens	19391			<u> </u>				i
Brevibacterium	linens	8377							
Brevibacterium	paraffinolyticum					11160			
Brevibacterium	spec.				1		717.73		<u> </u>
Brevibacterium	spec.						717.73		
Brevibacterium	spec.	14604							
Brevibacterium	spec.	21860							
Brevibacterium	spec.	21864	<u> </u>					 	
Brevibacterium	spec.	21865							
Brevibacterium	spec.	21866							
Brevibacterium	spec.	19240							
Corynebacterium	acetoacidophilum	21476							
Corynebacterium	acetoacidophilum	13870							
Corynebacterium	acetoglutamicum			B11473					
Corynebacterium	acetoglutamicum			B11475					
Corynebacterium	acetoglutamicum	15806		-					
Corynebacterium	acetoglutamicum	21491						i	
Corynebacterium	acetoglutamicum	31270							
Corynebacterium	acetophilum			B3671					
Corynebacterium	ammoniagenes	6872					-	2399	
Corynebacterium	ammoniagenes	15511						····	
Corynebacterium	fujiokense	21496							
Corynebacterium	glutamicum	14067							
Corynebacterium	glutamicum	39137				******			
Corynebacterium	glutamicum	21254	i				*		
Corynebacterium	glutamicum	21255							••
Corynebacterium	glutamicum	31830							
Corynebacterium	glutamicum	13032							
Corynebacterium	glutamicum	14305							
Corynebacterium	glutamicum	15455							
Corynebacterium	glutamicum	13058							
Corynebacterium	glutamicum	13059							
Corynebacterium	glutamicum	13060							
Corynebacterium	glutamicum	21492							
Corynebacterium	glutamicum	21513							
Corynebacterium	glutamicum	21526					-		
Corynebacterium	glutamicum	21543							
Corynebacterium	glutamicum	13287							
Corynebacterium	glutamicum	21851							
Corynebacterium	glutamicum	21253				-			
Corynebacterium	glutamicum	21514							
	glutamicum	21516							
	glutamicum	21299							

Genus 🔭 🥞	species	ATCC	FERM.	NRRL	CECT	NŒIMB	CBS	NCTC	DSMZ
Corynebacterium	glutamicum	21300							
Corynebacterium	glutamicum	39684			1				
Corynebacterium	glutamicum	21488			1				
Corynebacterium	glutamicum	21649							
Corynebacterium	glutamicum	21650			ļ			<u> </u>	
Corynebacterium	glutamicum	19223			 			 	
Corynebacterium	glutamicum	13869	l					_	
Corynebacterium	glutamicum	21157							
Corynebacterium	glutamicum	21158	-						
Corynebacterium	glutamicum	21159	-						
Corynebacterium	glutamicum	21355							
Corynebacterium	glutamicum	31808							
Corynebacterium	glutamicum	21674			<u> </u>				-
Corynebacterium	glutamicum	21562							
Corynebacterium	glutamicum	21563	-						
Corynebacterium	glutamicum	21564							
	glutamicum	21565							
•	glutamicum	21566							
Corynebacterium	glutamicum	21567							
	glutamicum	21568							
	glutamicum	21569							
<u> </u>	glutamicum	21570							
	glutamicum	21571							
<u> </u>	glutamicum	21572							-
Corynebacterium	glutamicum	21573							
	glutamicum	21579				-			
	glutamicum	19049							
	glutamicum	19050							
Corynebacterium	glutamicum	19051							
Corynebacterium	glutamicum	19052							
Corynebacterium	glutamicum	19053							
Corynebacterium	glutamicum	19054							
Corynebacterium	glutamicum	19055				i			
Corynebacterium	glutamicum	19056							
Corynebacterium	glutamicum	19057							
Corynebacterium	glutamicum	19058							
Corynebacterium	glutamicum	19059							
Corynebacterium	glutamicum	19060					,		
Corynebacterium	glutamicum	19185							
Corynebacterium	glutamicum	13286							
Corynebacterium	glutamicum	21515							
	glutamicum	21527							
	glutamicum	21544			1				
	glutamicum	21492		i	- 1				
	glutamicum			B8183					
	glutamicum			B8182					
	glutamicum			B12416					
	glutamicum			B12417					

Genus :	species	#ATCC	FERM	NRRE	CECT	NCIMB	CBS	NCTO	DSMZ
Corynebacterium	glutamicum			B12418				Service Services	
Corynebacterium	glutamicum	_		B11476					
Corynebacterium	glutamicum	21608							
Corynebacterium	lilium		P973			-			
Corynebacterium	nitrilophilus	21419				11594			
Corynebacterium	spec.		P4445						
Corynebacterium	spec.		P4446						
Corynebacterium	spec.	31088							-
Corynebacterium	spec.	31089							
Corynebacterium	spec.	31090							
Corynebacterium	spec.	31090	-						
Corynebacterium	spec.	31090							
Corynebacterium	spec.	15954							20145
Corynebacterium	spec.	21857	_						
Corynebacterium	spec.	21862							
Corynebacterium	spec.	21863							

ATCC: American Type Culture Collection, Rockville, MD, USA

FERM: Fermentation Research Institute, Chiba, Japan

NRRL: ARS Culture Collection, Northern Regional Research Laboratory, Peoria, IL, USA

CECT: Coleccion Espanola de Cultivos Tipo, Valencia, Spain

NCIMB: National Collection of Industrial and Marine Bacteria Ltd., Aberdeen, UK

CBS: Centraalbureau voor Schimmelcultures, Baarn, NL NCTC: National Collection of Type Cultures, London, UK

DSMZ: Deutsche Sammlung von Mikroorganismen und Zellkulturen, Braunschweig, Germany

For reference see Sugawara, H. et al. (1993) World directory of collections of cultures of microorganisms: Bacteria, fungi and yeasts (4th edn), World federation for culture collections world data center on microorganisms, Saimata, Japen.

WO 01	/0084	14					!	90				PCT/I	B 00/009	43	
<u>Date of</u> Deposit	13-Jul-99	20-Sep-99	20-Sep-99	17-Jun-98	8-Aug-97	28-Feb-96 20-Apr-98	11-MAR-1999	12-Jul-99	12-MAY-1998	22-OCT-1997	16-Jul-98	08-OCT-1997 (Rel. 52,	07-OCT-1996 21-OCT-1999	08-OCT-1997 (Rel. 52,	07-OCT-1996
% homology. Date of (GAP)	37,148	34,568	34,568	58,140	57,589	55,667 45,283	42,991	44,444	39,689	48,045	38,514	99,031	99,031 43,663	94,767	94,767
Source of Genbank Hit	Homo sapiens	Drosophila melanogaster	Drosophila melanogaster	Mycobacterium	tuberculosis Mycobacterium leprae	Streptomyces anulatus n Dictyostelium discoideum	Danio rerio	n Dictyostelium discoideum	Rhodobacter capsulatus	g Rhodobacter sphaeroides	Klebsiella pneumoniae	Corynebacterium glutamicum	Unknown. . Leishmania major	Corynebacterium glutamicum	Unknown.
Table 4: Alignment Results Length Accession Name of Genbank Hit	HS_5402_B2_A12_T7A RPCI-11 Human Male BAC Library Homo sapiens denomic clone Plate=978 Col=24 Row=B. denomic survey sequence	Drosophila melanogaster chromosome 2 clone BACR07M10 (D630) RPCI-98 07.M.10 map 24A-24D strain y; cn bw sp, *** SEQUENCING IN PROGRESS ***, 83 unordered pieces.	Drosophila melanogaster chromosome 2 clone BACR07M10 (D630) RPCI-98 07.M.10 map 24A-24D strain y; cn bw sp, *** SEQUENCING IN PROGRESS***, 83 unordered pieces.	Mycobacterium tuberculosis H37Rv complete genome; segment 136/162.	Mycobacterium leprae cosmid B1779.	S.alboniger napH, pur7, pur10, pur6, pur4, pur5 and pur3 genes. C89713 Dictyostellum discoideum SS (H. Urushihara) Dictyostellum discoideum discoideum	cDNA clone SSG229, mRNA sequence. fb63g03.y1 Zebrafish WashU MPIMG EST Danio rerio cDNA 5' similar to SW:AFP4_MYOOC P80961 ANTIFREEZE PROTEIN LS-12. ;; mRNA	sequence. C92167 Dictyostelium discoideum SS (H.Urushihara) Dictyostelium discoideum Dictyostelium discoideum cDNA clone SSD179, mRNA sequence.	Rhodobacter capsulatus strain SB1003, partial genome.	Rhodobacter sphaeroides operon regulator (smoC), periplasmic sorbitol-binding Rhodobacter sphaeroides protein (smoE), sorbitol/mannitol transport inner membrane protein (smoF), sorbitol/mannitol transport inner membrane protein (smoG), sorbitol/mannitol transport inner membrane protein (smoG), sorbitol/mannitol transport protein (smoK), sorbitol dehydrogenase (smoS), mannitol dehydrogenase (mtlK), and periplasmic mannitol-binding protein (smoM) genes, complete cds.	Klebsiella pneumoniae D-arabinitol transporter (daIT), D-arabinitol kinase (daIK), D-arabinitol dehydrogenase (daID), and repressor (daIR) genes, complete cds.	Base sequence of sucrase gene.	Sequence 4 from patent US 5556776. Leishmania major Friedlin chromosome 23 cosmid L5883, complete sequence.	Base sequence of sucrase gene.	Sequence 4 from patent US 5556776.
Accession	AQ713475	130583 AC007420	130583 AC007420	Z83867	Z98271	X92429 C89713	Al497294	C92167	189370 AF010496	AF018073	AF045245	E11760	126124 AL117384	E11760	126124
Length	581	130583	130583	25830	43254	9120 767	484	637	189370	9810	5930	6911	6911 31934	6911	6911
length. Genbank Hit (NT)	GB_GSS4:AQ713475	GB_HTG3:AC007420	GB_HTG3:AC007420	GB_BA1:MTCY3A2	GB_BA1:MLCB1779	GB_BA1:SAPURCLUS GB_EST21:C89713	GB_EST28:AI497294	GB_EST21:C92167	GB_BA2:AF010496	GB_BA2:AF018073	GB_BA2:AF045245	ка00041 1342. EM_PAT.E11760	GB_PAT:126124 GB_IN1:LMFL5883	EM_PAT:E11760	GB_PAT:126124
ID# length (NT)	rxa00013 996			rxa00014 903		гха00030 513			rxa00032 1632			rxa00041 1342		rxa00042 882	

<i>1</i> 0.5							· •						•	C1/1100	70024	.5
23-Jan-96 07-OCT-1996 08-OCT-1997 (Rel. 52,	Created) 24-Jun-98 19-Apr-97	27-Jul-98 17-Jun-98	17-Jun-98	03-DEC-1996	10-DEC-1996	03-DEC-1996	10-DEC-1996	17-Jun-98	19-Jun-98	15-Jun-96 19-Jun-98	19-Jun-98	19-Jun-98	27-Apr-93 13-MAR-1997	2-Feb-99	2-Feb-99	17-Sep-97 23-Jun-99
40,276 97,591 97,591	35,879 is 62,658	37,638 36,784	67,457	40,883	67,457	35,883	51,001	51,001	35,735	57,014 41,892	41,841	36,599	36,212 38,816	42,239	37,307	58,312 36,632
Caenorhabditis elegans Únknown. Corynebacterium glutamicum	Homo sapiens Mycobacterium smegmatis 62,658	Streptomyces coelicolor Mycobacterium	tuberculosis Mycobacterium	tuberculosis Mycobacterium	tuberculosis Mycobacterium	tuberculosis Mycobacterium	tuberculosis Mycobacterium	tuberculosis Mycobacterium	tuberculosis Mycobacterium	tuberculosis Mycobacterium leprae Mycobacterium	tuberculosis Mycobacterium	tuberculosis Mycobacterium	tuberculosis Rattus norvegicus ', Mus musculus	'Mus musculus	Mus musculus	Mycobacterium leprae Mycobacterium tuberculosis
Table 4 (continued) Caenorhabditis elegans sur-2 mRNA, complete cds. Sequence 4 from patent US 5556776. Base sequence of sucrase gene.	 Homo sapiens clone UWGC:g1564a012 from 7p14-15, complete sequence. Mycobacterium smegmatis phosphoglucose isomerase gene, complete cds. 	Streptomyces coelicolor cosmid 5A7. Mycobacterium tuberculosis H37Rv complete genome; segment 44/162.	Mycobacterium tuberculosis H37Rv complete genome; segment 65/162.	Mycobacterium tuberculosis sequence from clone y456.	Mycobacterium tuberculosis sequence from clone y175.	Mycobacterium tuberculosis sequence from clone y456.	Mycobacterium tuberculosis sequence from clone y175.	Mycobacterium tuberculosis H37Rv complete genome; segment 65/162.	Mycobacterium tuberculosis H37Rv complete genome; segment 126/162.	Mycobacterium leprae cosmid B1529 DNA sequence. Mycobacterium tuberculosis H37Rv complete genome; segment 126/162.	Mycobacterium tuberculosis H37Rv complete genome; segment 126/162.	Mycobacterium tuberculosis H37Rv complete genome; segment 126/162.		mRNA sequence. mw96a03.y1 Soares mouse NML Mus musculus cDNA clone IMAGE:678508 5' Mus musculus similar to TR:009171 009171 BETAINE-HOMOCYSTEINE	mw95c10.y1 Soares mouse NML Mus musculus cDNA clone IMAGE:678450 51 mRNA sequence	Mycobacterium leprae cosmid B637. Mycobacterium tuberculosis H37Rv complete genome; segment 132/162.
U33051 26124 E11760	AC005174 U88433	AL031107 Z79700	279701	AD000001	AD000015	AD000001	AD000015	279701	Z74024	L78824 Z74024	Z74024	Z74024	M55532 AA253618	Al390284	AI390280	Z99263 AL021287
4899 6911 6911	39769 1928	40337 39800	38300	37316	18106	37316	18106	38300	39991	39991 39991	39991	39991	10752 313	490	467	44882 70287
GB_IN1:CEU33051 GB_PAT:I26124 EM_PAT:E11760	GB_PR3:AC005174 GB_BA1:MSU88433	GB_BA1:SC5A7 GB_BA1:MTCY10D7	GB_BA1:MTCY277	GB_BA1:MSGY456	GB_BA1:MSGY175	GB_BA1:MSGY456	GB_BA1:MSGY175	GB_BA1:MTCY277	GB_BA1:MTCY274	GB_BA1:MSGB1529CS 36985 GB_BA1:MTCY274 39991	GB_BA1:MTCY274	GB_BA1:MTCY274	GB_RO:RATCBRQ GB_EST11:AA253618	GB_EST26:Al390284	GB_EST26:Al390280	GB_BA1:MLCB637 GB_BA1:MTV012
ка00043 1287	rxa00098 1743		rxa00148 2334			rxa00149 1971			rxa00195 684		xa00196 738		xa00202 1065			rxa00206 1161

PCT/IB00/00943

WO 01/00844

				Table 4 (continued)			
	GB_BA1:SC6E10	23990	AL109661	Streptomyces coelicolor cosmid 6E10.	Streptomyces coelicolor	38,616	5-Aug-99
xa00224 1074	GB_BA1:BJU32230	1769	U32230	A3(2) Bradyrhizobium japonicum electron transfer flavoprotein small subunit (etfS) nd Bradyrhizobium japonicum 48,038	A3(2) Bradyrhizobium japonicum	48,038	25-MAY-1996
	GB_BA1:PDEETFAB	2440	L14864	large subunit (etfL) genes, complete cds. Paracoccus denitrificans electron transfer flavoprotein alpha and beta subunit	Paracoccus denitrificans	48,351	27-OCT-1993
	GB HTG3:AC009689	177954	177954 AC009689	genes, complete cds's. Homo saciens chromosome 4 clone 104 F 7 map 4 1 OW-PASS SEQUENCE Homo saciens	Homo sapiens	38 756	28-Aug-99
				SAMPLING.		3	56-55V-07
xa00225 909	GB_RO:AF060178	2057	AF060178	Mus musculus heparan sulfate 2-sulfotransferase (Hs2st) mRNA, complete cds. Mus musculus	.Mus musculus	39,506	18-Jun-98
٠	GB_GSS11:AQ325043	734	AQ325043	mgxb0020J01r CUGI Rice Blast BAC Library Magnaporthe grisea genomic	Magnaporthe grisea	38,333	8-Jan-99
	GB_EST31:AI676413	551	AI676413	clone mgxb0020J01r, genomic survey sequence. etmEST0167 EtH1 Eimeria tenella cDNA clone etmc074 5', mRNA sequence.	Eimeria tenella	35,542	19-MAY-1999
xa00235 1398	GB_BA1:MTCY10G2	38970	Z92539	Mycobacterium tuberculosis H37Rv complete genome; segment 47/162.	Mycobacterium	65,759	17-Jun-98
	GB_BA2:AF061753	3721	AF061753	Nitrosomonas europaea CTP synthase (pyrG) gene, partial cds; and enolase	Nitrosomonas europaea	58,941	31-Aug-98
	GB_BA2:AF086791	37867	AF086791	Ymory gons, comprose 25. Ymomonas mobilis strain ZM4 clone 67E10 carbamoylphosphate synthetase small subunit (carA), carbamoylphosphate synthetase large subunit (carB), transcription elongation factor (greA), enolase (eno), pyruvate dehydrogenase alpha subunit (pdhA), pyruvate dehydrogenase beta subunit (pdhB), ribonuclease H (rnh), homoserine kinase homolog, alcohol dehydrogenase II	Zymomonas mobilis	61,239	4-Nov-98
				(adhB), and excinuclease ABC subunit A (uvrA) genes, complete cds; and unknown genes.			
rxa00246 1158	GB_BA2:AF012550	2690	AF012550	Acinetobacter sp. BD413 ComP (comP) gene, complete cds.	Acinetobacter sp. BD413	53,726	27-Sep-99
	GB_PAT:E03856	1506	E03856	gDNA encoding alcohol dehydrogenase.	Bacillus	51,688	29-Sep-97
	GB_BA1:BACADHT	1688	D90421	B.stearothermophilus adhT gene for alcohol dehydrogenase.	stearothermophilus Bacillus	51,602	7-Feb-99
rxa00251 831	GB_BA1:MTCY20G9	37218	Z77162	Mycobacterium tuberculosis H37Rv complete genome; segment 25/162.	stearothermophilus Mycobacterium	42,875	17-Jun-98
	GB_BA1:MTV004	69350	AL009198	Mycobacterium tuberculosis H37Rv complete genome; segment 144/162.	tuberculosis Mycobacterium	40,380	18-Jun-98
	GB_BA1:MTV004	69350	AL009198	Mycobacterium tuberculosis H37Rv complete genome; segment 144/162.	tuberculosis Mycobacterium tuberculosis	41,789	18-Jun-98
rxa00288 1134	GB_BA2:AF050114	1038	AF050114	Pseudomonas sp. W7 alginate lyase gene, complete cds.	Pseudomonas sp. W7	49,898	03-MAR-1999
	GB_GSS3:B16984	469	B16984	344A14 TVC CIT978SKA1 Homo sapiens genomic clone A-344A14, genomic survey sequence.	Homo sapiens	39,355	4-Jun-98
ma00293 1035	GB_IN2:AF144549 GB_EST1:T28483	7887 313	AF144549 T28483	Aedes albopictus ribosomal protein L34 (rpl34) gene, complete cds. EST46182 Human Kidney Homo sapiens cDNA 3' end similar to flavin-containing monooxygenase 1 (HT:1956), mRNA sequence.	Aedes albopictus Homo sapiens	36,509 42,997	3-Jun-99 6-Sep-95

Table 4 (continued)				
Human flavin-containing monooxygenase (FMO1) mRNA, complete cds.	Homo sapiens	37,915	8-Nov-94	W
Zbr. 3c03.y3 Soares_retal_lung_NbHL19W Homo sapiens cDNA clone IMAGE:309224 5' similar to gb:M64082 DIMETHYLANILINE MONOOXYGENASE (HUMAN): mRNA sequence.	Homo sapiens	41,502	14-Jun-99	O 01/0
9 Drosophila melanogaster chromosome X clone BACR11H20 (D881) RPCI-98 11.H.20 map 12B-12C strain y; cn bw sp, *** SEQUENCING IN PROGRESS	Drosophila melanogaster	33,890	02-DEC-1999	0844
 ijosa i z.s.i NCI_CGAP_Prio Homo sapiens cDNA clone IMAGE:997175, mRNA sequence. 	Homo sapiens	40,821	20-Aug-97	
Drosophila melanogaster chromosome X clone BACR11H20 (D881) RPCI-98 11.H.20 map 12B-12C strain y; cn bw sp, *** SEQUENCING IN PROGRESS ***, 92 unordered pieces.	Drosophila melanogaster	30,963	02-DEC-1999	
variola minor virus complete genome		6		
Variola maior virus (strain Bandladesh-1975) complete genome	Variola minor virus	35,883	2-Sep-99	
Variola virus DNA complete genome	Variora major virus	34,664	12-Jan-95	
Homo sapiens chromosome 4 clone 57 A 22 map 4 *** SEQUENCING IN	Variota virus Homo espisos	36,000	13-DEC-1996	
PROGRESS ***, 8 unordered pieces.	significance of the second of	30,988	29-Sep-99	
Homo sapiens chromosome 4 clone 57_A_22 map 4, *** SEQUENCING IN PROGRESS *** 8 unordered places	Homo sapiens	36,988	29-Sep-99	
Homo sapiens chromosome 17. clone hRPK 138 P. 22. complete segments				9
L.casei gene for ATPase beta-subunit.	nomo sapiens Lactobacillus pasai	36,340	09-OCT-1998	3
L.casei gene for ATPase beta-subunit.	Lactobacillus casei	39,308	11-DEC-1992 11-DEC-1992	
Salmonella (S2980) proline nemesso (Ottus) sacamen		;	;	
commenced (promise permease (pure) gene, 3 end.	Salmonella sp.	39,623	09-MAY-1996	
Salmonella (S2983) proline permease (putP) gene, 5' end.	Salmonella sp.	39,623	09-MAY-1996	
Salmonella (S3015) proline permease (putP) gene, 5' end.	Salmonella sp.	42,906	09-MAY-1996	
Homo sapiens PAC clone DJ0740D02 from 7p14-p15, complete sequence.	Homo sapiens	38,142	16-MAY-1998	
Homo sapiens clone DJ0891L14, complete sequence.	Homo sapiens	38.549	17-Jul-99	
Homo sapiens PAC clone DJ0740D02 from 7p14-p15, complete sequence.	Homo sapiens	35,865	16-MAY-1998	
Mycobacterium tuberculosis H37Rv complete genome; segment 99/162.	Mycobacterium	38,940	24-Jun-99	PC T/
RPCI-11-195H2.TV RPCI-11 Homo saniens genomic clone RPCI-11-105U2	tuberculosis			тв
genomic survey sequence.	supplements	30'33	23-MAK-1999	00/0
Astasia longa small subunit ribosomal RNA gene, complete sequence.	Astasia longa	36.465	28-Jun-99	909
Caenorhabditis elegans chromosome III clone Y56A3, *** SEQUENCING IN	Caenorhabditis elegans	35,179	6-Sep-99) 43
Caenorabditis elegans chromosome III clone Y56A3, *** SEQUENCING IN	Caenorhabditis elegans	35,179	6-Sep-99	
PROGRESS ***, in unordered pieces.			22	

224746 AL022280

GB_HTG1:CEY56A3

AQ412290

238

GB_GSS12:AQ412290

Z70692

38110

GB_BA1:MTCY427

xa00340 1269

AF112871 AL022280

2394

GB_PL2:AF112871 GB_HTG1:CEY56A3

rxa00379 307

224746

129014 AC004916

GB_PR4:AC004916 GB_PR3:AC004691

GB_PR3:AC004691

141990 AC004691

141990 AC004691

rxa00329 1347

1889

GB_BA1:STYPUTPI

159648 AC009571 159648 AC009571

GB_HTG3:AC009571 GB_HTG3:AC009571

AC005697

174503

X64542

1514 1514

GB_BA1:LCATPASEB GB_BA1:LCATPASEB GB_BA1:LCATPASEB

xa00327 507

X64542

L01138

1887

GB_BA1:STYPUTPE GB_BA1:STYPUTPF

rxa00328 615

L01139 L01142

1887

186986 Y16780 185578 X69198 186103 L22579

GB_VI:VMVY16780

xa00310 558

GB_VI:WCGAA GB_VI:VARCG

rxa00317 777

AI734238

M64082

2134 512

GB_PR1:HUMFM01 GB_EST32:AI734238

168266 AC011069

GB_HTG6:AC011069

xa00296 2967

AA531468

GB_EST15:AA531468 414

168266 AC011069

GB_HTG6:AC011069

	GB_PR2:HS134019	86897	AL034555	Table 4 (continued) Human DNA sequence from clone 134019 on chromosome 1p36.11-36.33,	Homo sapiens	40,604	23-Nov-99
rxa00381 729	GB_GSS4:AQ730532	416	AQ730532	complete sequence. HS_2149_A1_C06_T7C CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2149 Col=11 Row=E, genomic survey sequence.	Homo sapiens	35,766	15-Jul-99
	GB_EST23:AI120939	561	AI120939	ub74f05.r1 Soares mouse mammary gland NMLMG Mus musculus cDNA clone Mus musculus IMAGE:1383489 5' similar to gb:J04046 CALMODULIN (HUMAN); gb:M19381 Mouse calmodulin (MOUSE): mRNA sequence.	Mus musculus	41,113	2-Sep-98
	GB_EST23:AI120939	561	Al120939	ub74f05.r1 Soares mouse mammary gland NMLMG Mus musculus cDNA clone Mus musculus IMAGE:1383489 5' similar to gb:J04046 CALMODULIN (HUMAN); gb:M19381 Mouse calmodulin (MOUSE);, mRNA sequence.	Mus musculus	41,113	2-Sep-98
rxa00385 362	GB_EST32:AI726450	565	AI726450	BNLGHi5857 Six-day Cotton fiber Gossypium hirsutum cDNA 5' similar to (AF015913) Skb1Hs [Homo sapiens], mRNA sequence.	Gossypium hirsutum	41,152	11-Jun-99
	GB_GSS4.AQ740856	768	AQ740856	HS_2274_A2_A07_T7C CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2274 Col=14 Row=A, genomic survey sequence.	Homo sapiens	41,360	16-Jul-99
	GB_PR1:HSPAIP	1587	X91809	H.sapiens mRNA for GAIP protein.	Homo sapiens	36,792	29-MAR-1996
rxa00388 1134	GB_BA1:MTY25D10	40838	Z 95558	Mycobacterium tuberculosis H37Rv complete genome; segment 28/162.	Mycobacterium tuberculosis	51,852	17-Jun-98
	GB_BA1:MSGY224	40051	AD000004	Mycobacterium tuberculosis sequence from clone y224.	Mycobacterium tuberculosis	51,852	03-DEC-1996
	GB_HTG1:AP000471	72466	AP000471	Homo sapiens chromosome 21 clone B2308H15 map 21q22.3, *** SEQUENCING IN PROGRESS ***, in unordered pieces.	Homo sapiens	36,875	13-Sep-99
rxa00427 909	GB_BA1:MSGY126	37164	AD000012	Mycobacterium tuberculosis sequence from clone y126.	Mycobacterium	60,022	10-DEC-1996
	GB_BA1:MTY13D12	37085	Z80343	Mycobacterium tuberculosis H37Rv complete genome; segment 156/162.	tuberculosis Mycobacterium tuberculosis	60,022	17-Jun-98
	GB_HTG1:CEY48C3	270193	292855	Caenorhabditis elegans chromosome II clone Y48C3, *** SEQUENCING IN PROGRESS *** in unordered pieces.	Caenorhabditis elegans	28,013	29-MAY-1999
xa00483 1587	GB_PR2:HSAF001550		173882 AF001550	Homo sapiens chromosome 16 BAC clone CIT987SK-334D11 complete	Homo sapiens	38,226	22-Aug-97
	GB_BA1:LLCPJW565	12828	Y12736	Lactococcus lactis cremoris plasmid pJW565 DNA, abiiM, abiiR genes and	Lactococcus lactis subsp.	37,492	01-MAR-1999
	GB_HTG2:AC006754	206217	206217 AC006754	Caenorhabditis elegans clone Y40B10, *** SEQUENCING IN PROGRESS ***, 5 unordered nieces	cremoris Caenorhabditis elegans	36,648	23-Feb-99
rxa00511 615	GB_PR3:HSE127C11	38423	Z 74581	Human DNA sequence from cosmid E127C11 on chromosome 22q11.2-qter contains STS.	Homo sapiens	39,831	23-Nov-99
	GB_PR3:HSE127C11	38423	Z74581	Human DNA sequence from cosmid E127C11 on chromosome 22q11.2-qter contains STS.	Homo sapiens	36,409	23-Nov-99
rxa00512 718	GB_BA1:MTCY22G8	22550	295585	Mycobacterium tuberculosis H37Rv complete genome; segment 49/162.	Mycobacterium tuberculosis	56,232	17-Jun-98

PCT/IB00/00943

WO 01/00844

28-Sep-99

37,143

37,976

38,732

43481 AB026648 Arabidopsis thaliana genomic DNA, chromosome 3, P1 clone: MLJ15, complete Arabidopsis thaliana

sedneuce.

Homo sapiens chromosome 19 clone CITB-E1_2568A17, *** SEQUENCING IN Homo sapiens PROGRESS *** 40 unordered pieces.

Homo sapiens chromosome 19 clone CiTB-E1_2568A17, *** SEQUENCING IN Homo sapiens PROGRESS ***, 40 unordered pieces.

181745 AC008179 Homo sapiens clone NH0576F01, complete sequence.

197110 AC010325

GB_HTG3:AC010325

GB_PR4:AC008179

197110 AC010325

GB_HTG3:AC010325

rxa00682 2022

GB_PL1:AB026648

37,976

				Table 4 (continued)		
	GB_BA1:MSGLTA	1776	X60513	M.smegmatis gltA gene for citrate synthase.	Mycobacterium smegmatis 56,143	56,143
	GB_BA2:ECU73857	128824	U73857	Escherichia coli chromosome minutes 6-8.	Escherichia coli	48,563
rxa00517 1164	GB_HTG2:AC006911	298804	AC006911	AC006911 Caenorhabditis elegans clone Y94H6x, *** SEQUENCING IN PROGRESS ***, Caenorhabditis elegans	Caenorhabditis elegans	37,889
				15 unordered pieces.		
	GB_HTG2:AC006911	298804	AC006911	AC006911 Caenorhabditis elegans clone Y94H6x, *** SEQUENCING IN PROGRESS ***, Caenorhabditis elegans	Caenorhabditis elegans	37,889
				15 unordered pieces.		
	GB_EST29:AI602158 481	481	AI602158	UI-R-AB0-vy-a-01-0-UI.s2 UI-R-AB0 Rattus norvegicus cDNA clone UI-R-AB0- Rattus norvegicus	Rattus norvegicus	40,833
				vy-a-01-0-UI 3', mRNA sequence.	1	
xa00518 320	GB_BA2:ECU73857	128824	U73857	Escherichia coli chromosome minutes 6-8.	Escherichia coli	49,688
	GB_BA2:STU51879	8371	U51879	Salmonella typhimurium propionate catabolism operon: RpoN activator protein Salmonella typhimurium	Salmonella typhimurium	50,313

				homolog (prpR), carboxyphosphonoenolpyruvate phosphonomutase homolog (prpB), citrate synthase homolog (prpC), prpD and prpE genes,			
rxa00606 2378	GB_BA2:AE000140 GB_EST32:AU068253	12498 376	AE000140 AU068253	complete cds. Escherichia coli K-12 MG1655 section 30 of 400 of the complete genome. Escherichia « AU068253 Rice callus Oryza sativa cDNA clone C12658_9A, mRNA sequence. Oryza sativa	Escherichia coli Oryza sativa	49,688 41,333	12-Nov-98 7-Jun-99
	GB_EST13:AA363046 GB_EST32:AU068253	329 376	AA363046 AU068253	EST72922 Ovary II Homo sapiens cDNA 5' end, mRNA sequence. AU068253 Rice callus Oryza sativa cDNA clone C12658_9A, mRNA sequence. Oryza sativa	Homo sapiens Oryza sativa	34,347 41,899	21-Apr-97 7-Jun-99
rxa00635 1860	GB_BA1:PAORF1	1440	X13378	Pseudomonas amyloderamosa DNA for ORF 1.	Pseudomonas	53,912	14-Jul-95
	GB_BA1:PAORF1	1440	X13378	Pseudomonas amyloderamosa DNA for ORF 1.	amyloderamosa Pseudomonas amyloderamosa	54,422	14-Jul-95
rxa00679 1389	GB_PL2:AC010871	80381		l BAC T16011 genomic sequence,	Arabidopsis thaliana	38,244	13-Nov-99
	GB_PL1:AT81KBGEN	81493	X98130	complete sequence. A.thaliana 81kb genomic sequence.	Arabidopsis thaliana	36,091	12-MAR-1997
	GB_PL2:AC010871	80381		chromosome III BAC T16O11 genomic sequence,	Arabidopsis thaliana	37,135	13-Nov-99
rxa00680 441	GB_PR3:AC004058	38400	AC004058	complete sequence. 38400 AC004058 Homo sapiens chromosome 4 clone B241P19 map 4q25, complete sequence.	Homo sapiens	36,165	30-Sep-98
	GB_PL1:AT81KBGEN 81493 X98130	81493	X98130	A.thaliana 81kb genomic sequence.	Arabidopsis thaliana	38,732	12-MAR-1997

				Table 4 (continued)			
rxa00683 1215	GB_BA2:AE000896	10701	AE000896	Methanobacterium thermoautotrophicum from bases 1189349 to 1200055		38,429	15-Nov-97
	GB IN1-DMBB7A4	212734	AI 109630	(section 102 of 146) of the complete genome. Drosophila melanopaster clone RACR7A4	mermoautotrophicum Drosnohila melanogaster	36 454	30-Jul-99
	5	273	AV163010	AV163010 Mus musculus head C57BL/6J 13-day embryo Mus musculus cDNA Mus musculus	0	41,758	8-Jul-89
гха00686 927	GB_HTG2:HSDJ137K2 190223	190223	AL049820	cione 3110005JZ2, mKNA sequence. Homo sapiens chromosome 6 clone RP1-137K2 map q25.1-25.3, *** SEQUENCING IN PROGRESS *** in unordered pieces.	Homo sapiens	38,031	03-DEC-1999
	GB_HTG2:HSDJ137K2 190223	190223	AL049820	Homo sapiens chromosome 6 clone RP1-137K2 map q25.1-25.3, ***	Homo sapiens	38,031	03-DEC-1999
	GB_EST12:AA284399	431	AA284399	zs57b04.r1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:701551 5',	Homo sapiens	39,205	14-Aug-97
rxa00700 927	GB_EST34:AI785570	454	AI785570	mRNA sequence. uj44d03.x1 Sugano mouse liver mlia Mus musculus cDNA clone	Mus musculus	41,943	2-Jul-99
				IMAGE:1922789 3' similar to gb:Z28407 60S RIBOSOMAL PROTEIN L8 (HUMAN):. mRNA sequence.			
	GB_EST25:Al256147	684	AI256147	UIGGE12.x1 Sugano mouse liver mila Mus musculus cDNA clone IMAGE:1890190 3' similar to gb:Z28407 60S RIBOSOMAL PROTEIN L8 (HUMAN): mRNA sequence.	Mus musculus	40,791	12-Nov-98
	GB_BA1:CARCG12	2079	X14979	C. aurantiacus reaction center genes 1 and 2.	Chloroflexus aurantiacus	37,721	23-Apr-91
xa00703 2409	GB_BA1:SC7H2	42655	AL109732	Streptomyces coelicolor cosmid 7H2.	Streptomyces coelicolor A3(2)	56,646	2-Aug-99
	GB_BA1:MTCY274	39991	Z74024	Mycobacterium tuberculosis H37Rv complete genome; segment 126/162.	Mycobacterium tuberculosis	37,369	19-Jun-98
	GB_BA2:REU60056	2520	N60056	Ralstonia eutropha formate dehydrogenase-like protein (cbbBc) gene, complete Ralstonia eutropha cds.	Ralstonia eutropha	51,087	16-OCT-1996
rxa00705 1038	GB_GSS15:AQ604477	505	AQ604477	HS_2116_B1_G07_MR CIT Approved Human Genomic Sperm Library D Homo Homo sapiens sapiens genomic clone Plate=2116 Col=13 Row=N, genomic survey sequence.	Homo sapiens	39,617	10-Jun-99
	GB_EST11:AA224340 443	443	AA224340	zr14e07.s1 Stratagene hNT neuron (#937233) Homo sapiens cDNA clone IMAGE:648804 3', mRNA sequence.	Homo sapiens	35,129	11-MAR-1998
	GB_EST5:N30648	291	N30648	yw77b02.s1 Soares_placenta_8to9weeks_2NbHP8to9W Homo sapiens cDNA clone IMAGE:258219 3', mRNA sequence.	Homo sapiens	43,986	5-Jan-96
rxa00782 1005	GB_BA1:MTCY10D7	39800	Z79700	Mycobacterium tuberculosis H37Rv complete genome; segment 44/162.	Mycobacterium tuberculosis	63,327	17-Jun-98
	GB_BA1:MLCL373	37304	AL035500	Mycobacterium leprae cosmid L373.	Mycobacterium leprae	62,300	27-Aug-99
	GB_BA2:AF128399	2842	AF128399	Pseudomonas aeruginosa succinyl-CoA synthetase beta subunit (sucC) and succinyl-CoA synthetase alpha subunit (sucD) genes, complete cds.	Pseudomonas aeruginosa	53,698	25-MAR-1999
rxa00783 1395	GB_HTG2:AC008158	118792	AC008158	Homo sapiens chromosome 17 clone hRPK.42_F_20 map 17, *** SEQUENCING IN PROGRESS ***, 14 unordered pieces.	Homo sapiens	35,135	28-Jul-99
	GB_HTG2:AC008158	118792	AC008158	Homo sapiens chromosome 17 clone hRPK.42_F_20 map 17, *** SEQUENCING IN PROGRESS ***, 14 unordered pieces.	Homo sapiens	35,135	28-Jul-99
	GB_PR3:AC005017	137176	137176 AC005017	Homo sapiens BAC clone GS214N13 from 7p14-p15, complete sequence.	Homo sapiens	35,864	8-Aug-98
rxa00794 1128	GB_BA1:MIV017	67200	AL021897	Mycobacterium tuberculosis H3/RV complete genome; segment 48/162.	Mycobacterium tuberculosis	40,551	24-Jun-99

WO 01/00844

	GB_BA1:MLCB1222 GB_PR2:HS151B14	34714 128942	AL049491 Z82188	Mycobacterium leprae cosmid B1222. Human DNA sequence from clone 151B14 on chromosome 22 Contains SOMATOSTATIN RECEPTOR TYPE 3 (SS3R) gene,pseudogene similar to ribosomal protein L39,RAC2 (RAS-RELATED C3 BOTULINUM TOXIN SUBSTRATE 2 (P21-RAC2)) gene ESTs, STSs, GSSs and CpG islands, complete sequence.	Mycobacterium leprae Homo sapiens	61,170 37,455.	27-Aug-99 16-Jun-99
rxa00799 1767	GB_PL2.AF016327 616 GB_HTG2:HSDJ319M7 128208	616	AF016327 AL079341	Hordeum vulgare Barperm1 (perm1) mRNA, partial cds. Homo sapiens chromosome 6 clone RP1-319M7 map p21.1-21.3, *** SEQUENCING IN PROGRESS *** in mordered pieces.	Hordeum vulgare Homo sapiens	41,311 36,845	01-OCT-1997 30-Nov-99
	GB_HTG2:HSDJ319M7 128208	128208	AL079341	Homo sapiens chromosome 6 clone RP1-319M7 map p21.1-21.3, *** SEQUENCING IN PROGRESS ***, in unordered pieces.	Homo sapiens	36,845	30-Nov-99
rxa00800 1227	GB_BA1:MTV022	13025	AL021925	Mycobacterium tuberculosis H37Rv complete genome; segment 100/162.	Mycobacterium tuberculosis	63,101	17-Jun-98
	GB_BA1:AB019513	4417	AB019513	Streptomyces coelicolor genes for alcohol dehydrogenase and ABC transporter, complete cds.	Streptomyces coelicolor	41,312	13-Nov-98
	GB_PL1:SCSFAARP	2008	X68020	S.cerevisiae SFA and ARP genes.	Saccharomyces cerevisiae 36,288	36,288	29-Nov-94
rxa00825 1056	GB_BA1:MTY15C10	33050	Z95436	Mycobacterium tuberculosis H37Rv complete genome; segment 154/162.	Mycobacterium tuberculosis	39,980	17-Jun-98
	GB_BA1:MLCB2548 GB_BA2:AF169031	38916 1141	AL023093 AF169031	Mycobacterium leprae cosmid B2548. Xanthomonas oryzae pv. oryzae putative sugar nucleotide	Mycobacterium leprae Xanthomonas oryzae pv.	39,435 46,232	27-Aug-99 14-Sep-99
rxa00871				epimerase/dehydratase gene, partial cds.	oryzae		
rxa00872 1077	GB_IN1:CEF23H12 GB_HTG2:AC007263	35564 167390	Z74472 AC007263	Caenorhabditis elegans cosmid F23H12, complete sequence. Homo sapiens chromosome 14 clone BAC 79J20 map 14q31, *** eFOLIENCING IM PROCEECE ***	Caenorhabditis elegans Homo sapiens	34,502 35,714	08-OCT-1999 24-MAY-1999
	GB_HTG2:AC007263	167390	167390 AC007263	SECUCING IN PROGRESS ***, 3 ordered pieces. Homo sapiens chromosome 14 clone BAC 79J20 map 14q31, *** SEQUENCING IN PROGRESS ***, 5 ordered pieces.	Homo sapiens	35,714	24-MAY-1999
rxa00879 2241	GB_BA1:MTV049	40360	AL022021	Mycobacterium tuberculosis H37Rv complete genome; segment 81/162.	Mycobacterium tuberculosis	36,981	19-Jun-98
	GB_PL2:CDU236897	1827	AJ236897	Candida dubliniensis ACT1 gene, exons 1-2.	Candida dubliniensis	38,716	1-Sep-99
xa00909 955	GB_PL1:CAACT1A GB_BA2:AF010496	3206 189370	X16377 AF010496	Candida albicans act1 gene for actin. Rhodobacter capsulatus strain SB1003, partial genome.	Candida albicans Rhodobacter capsulatus	36,610 51,586	10-Apr-93 12-MAY-1998
	GB_BA1:RMPHA	7888	X93358	Rhizobium meliloti pha[A,B,C,D,E,F,G] genes.	Sinorhizobium meliloti	48,367	12-MAR-1999
	GB_EST16:C23528	317	C23528	C23528 Japanese flounder spleen Paralichthys olivaceus cDNA clone HB5(2), mRNA sequence.	Paralichthys olivaceus	41,640	28-Sep-99
xa00913 2118	GB_HTG2:AC007734	188267	188267 AC007734	Homo sapiens chromosome 18 clone hRPK.44_O_1 map 18, *** SEQUENCING IN PROGRESS ***, 18 unordered pieces.	Homo sapiens	34,457	66-unr-9

				Table 4 (continued)			
	GB_HTG2:AC007734	188267	188267 AC007734	Homo sapiens chromosome 18 clone hRPK 44_O_1 map 18, *** SEQUENCING IN PROGRESS ***, 18 unordered pieces.	Homo sapiens	34,457	66-unr-9
	GB_EST18:AA709478	406	AA709478	vv34a05.r1 Stratagene mouse heart (#937316) Mus musculus cDNA clone IMAGE:1224272 5: mRNA sequence	Mus musculus	42,065	24-DEC-1997
rxa00945 1095	GB_HTG4:AC010351	220710	220710 AC010351	Homo sapiens chromosome 5 clone CITB-H1_2022B6, *** SEQUENCING IN PROGRESS ***, 68 unordered pieces.	Homo sapiens	36,448	31-OCT-1999
	GB_HTG4:AC010351	220710	220710 AC010351	Homo sapiens chromosome 5 clone CITB-H1_2022B6, *** SEQUENCING IN PROGRESS ***, 68 unordered pleces.	Homo sapiens	36,448	31-OCT-1999
	GB_BA1:MTCY05A6	38631	296072	Mycobacterium tuberculosis H37Rv complete genome; segment 120/162.	Mycobacterium tuberculosis	36,218	17-Jun-98
rxa00965							
rxa00999 1575	GB_PAT:E13660	1916	E13660	gDNA encoding 6-phosphogluconate dehydrogenase.	Corynebacterium	98,349	24-Jun-98
	GR RA1-MTCV350	36021	783850	Muschardorium tuhasanbaia U27Bu asamalata asasasa 19460	glutamicum	6	;
		2005	50003	Mycoodicenum tuberculosis no/rx complete genome; segment 64/162.	Mycobacterium tuberculosis	38,520	17-Jun-98
	GB_BA1:MLCB1788	39228	AL008609	Mycobacterium leprae cosmid B1788.	Mycobacterium leprae	64,355	27-Aug-99
xa01015 442	GB_BA1:MIV008	63033	AL021246	Mycobacterium tuberculosis H37Rv complete genome; segment 108/162.	Mycobacterium	39,860	17-Jun-98
	GB_BA1:MTV008	63033	AL021246	Mycobacterium tuberculosis H37Rv complete genome; segment 108/162.	Mycobacterium	39,120	17-Jun-98
					tuberculosis		
xa01025 1119	GB_BA1:SC7A1 32039	32039	AL034447	Streptomyces coelicolor cosmid 7A1.	Streptomyces coelicolor	55,287	15-DEC-1998
	GB_BA1:MSGB1723C	S 38477	L78825	3 DNA sequence.	Mycobacterium leprae	56,847	15-Jun-96
	GB_BA1:MLCB637	44882	Z99263	Mycobacterium leprae cosmid B637.	Mycobacterium leprae	56,676	17-Sep-97
rxa01048 1347	GB_BA2:AF017444	3067	AF017444	Sinorhizobium melitoti NADP-dependent malic enzyme (tme) gene, complete cds.	Sinorhizobium meliloti	53,660	2-Nov-97
	GB_BA1:BSUB0013	218470	218470 Z99116	Bacillus subtilis complete genome (section 13 of 21): from 2395261 to 2613730	Bacillus subtilis	37,255	26-Nov-97
	GB_VI:HSV2HG52	154746	154746 Z86099	Herpes simplex virus type 2 (strain HG52), complete genome.	human herpesvirus 2	38.081	04-DEC-1998
rxa01049 1605	GB_HTG2:AC002518	131855	AC002518	Homo sapiens chromosome X clone bWXD20, *** SEQUENCING IN	Homo sapiens	35,647	2-Sep-97
			,	PROGRESS ***, 11 unordered pieces.			•
	GB_HTG2:AC002518	131855	131855 AC002518	Homo sapiens chromosome X clone bWXD20, *** SEQUENCING IN	Homo sapiens	35,647	2-Sep-97
	GB LTC2:ACOCCE48	124055	1210EE ACOOSE10	PROGRESS ***, 11 unordered pieces.		;	!
		2	20000	PROGRESS *** 11 unordered pieces.	one sapiens	70,100	/s-dac->
rxa01077 1494	GB_PR3:HSDJ653C5	85237	AL049743	Human DNA sequence from clone 653C5 on chromosome 1p21.3-22.3	Homo sapiens	36.462	23-Nov-99
				Contains CA repeat(D1S435), STSs and GSSs, complete sequence.	-		
	GB_BA1:ECU29579	72221	U29579	Escherichia coli K-12 genome; approximately 61 to 62 minutes.	Escherichia coli	41,808	1-Jul-95
04000	GB_BA1:ECU29579	72221	U29579	Escherichia coli K-12 genome; approximately 61 to 62 minutes.	Escherichia coli	36,130	1-Jul-95
radioes 673	GB_G336;AQU44021	38/	AQ044021	CII-HSP-2318C18.1R CII-HSP Homo sapiens genomic clone 2318C18, genomic survey sequence.	Homo sapiens	36,528	14-Jul-98

	GB_GSS8:AQ042907	392	AQ042907	CIT-HSF	Homo sapiens	35,969	14-Jul-98	•••
	GB_GSS8:AQ044021	387	AQ044021	genomic survey sequence. CIT-HSP-2318C18.TR CIT-HSP Homo sapiens genomic clone 2318C18.	Homo sapiens	44 545	14. hid-98	
rxa01093 1554	GB_BA1:CORPYKI	2795	L27126	genomic survey sequence. Corynebacterium pyruvate kinase gene, complete cds.	Corynebacterium	100 000	07-DEC-1004	, 500
	GB BA1:MTCY01B2	35938	295554	Woohactarium tuharrulosis H37Dv complete concessions and 201600	glutamicum			
				mycoogean in the control of the complete genome, segment 72/102.	Mycobacterium tuberculosis	63,771	17-Jun-98	
	GB_BA1:MIU65430	1439	U65430	Mycobacterium intracellulare pyruvate kinase (pykF) gene, complete cds.	Mycobacterium	67,061	23-DEC-1996	
rxa01099 948	GB_BA2:AF045998	780	AF045998	Corynebacterium glutamicum inositol monophosphate phosphatase (impA)	intracellulare Coronebacterium	90 616	10 00	
				gene, complete cds.	glutamicum		06-09-1-61	
	GB_BA2:AF051846	738	AF051846	Corynebacterium glutamicum phosphoribosylformimino-5-amino-1- phosphoribosyl-4- imidazolecarboxamide isomerase (hisA) gene, complete		100,000	12-MAR-1998	
	GB_GSS1:FR0005503	619	Z89313	cds. F.rubripes GSS sequence, clone 079B16aE8, genomic survey sequence.	Fugu rubripes	37,785	01-MAR-1997	
ma01111 541	GB_PR3:AC004063 GB_PR3:HS1178I21	177014 62268	177014 AC004063 62268 AL109852	Homo sapiens chromosome 4 clone B3218, complete sequence. Human DNA sequence from clone RP5-1178121 on chromosome X_complete.	Homo sapiens	35,835	10-Jul-98	
	TOSONO VICTORIA	0000	70000	sequence.	saplens	5/9/3	01-DEC-1999	
	GB_N165:AC009301	163369	163369 AC009301	Homo sapiens clone NH0062F14, *** SEQUENCING IN PROGRESS ***, 5 unordered pieces.	Homo sapiens	37,240	13-Aug-99	
rxa01130 687	GB_HTG3:AC009444	164587	164587 AC009444	Homo sapiens clone 1_O_3, *** SEQUENCING IN PROGRESS ***, 8	Homo sapiens	38,416	22-Aug-99	
	GB_HTG3:AC009444	164587	164587 AC009444	unordered pieces. Homo sapiens clone 1_O_3, *** SEQUENCING IN PROGRESS ***, 8	Homo sapiens	38,416	22-Aug-99	
rxa01193 1572	GB_IN1:DMC66A1	34127	AL031227 X76875	Disophia melangaster cosmid 66A1.	ogaster	38,416	05-OCT-1998	
		101	6 60 6	C.gradamicum (ASO 19) A I Pase Beta-subunit gene.	Corynebacterium	99,931	27-OCT-1994	
	EM_PAT:E09634	1452	E09634	Brevibacterium flavum UncD gene whose gene product is involved in	Corynebacterium	99,242	07-OCT-1997	
	•				glutamicum		(Rel. 52, Created)	
	GB_BA1:MLU15186	36241	U15186	Mycobacterium leprae cosmid L471.	Mycobacterium leprae	39,153	1995	
K801194 495	EM_PAT:E09634	1452	E09634	Brevibacterium flavum UncD gene whose gene product is involved in	Corynebacterium	100,000	1997	
					glutamicum		(Rel. 52,	TDA
	GB_BA1:CGASO19	1452	X76875	C.glutamicum (ASO 19) ATPase beta-subunit gene.	erium	100,000	1994	W/W
rxa01200	GB_VI:HEPCRE4B	414	X60570	Hepatitis C genomic RNA for putative envelope protein (RE4B isolate).	glutamicum Hepatitis C virus	36,769	5-Apr-92	1743

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01-MAY-1995	17-Jun-98	09-MAR-1995	01-MAY-1995	01-MAY-1995	26-MAY-1998	15-Sep-99	04-DEC-1999	04-DEC-1999	17-Jun-98	10-MAR-1998	26-Apr-93	2-Jun-98	17-Aug-99	17-Aug-99	13-MAR-1996	17-Jun-98	10-DEC-1996	09-OCT-1998 25-Sep-99	25-Sep-99
69,269	65,437	39,302	57,087	38,298	37,626	38,395	35,459	36,117	39,064	42,671	41,054	36,205	39,922	39,922	64,908	62,838	61,712	35,373 39,863	39,863
Streptomyces lividans	Mycobacterium tuberculosis	Mycobacterium leprae	Streptomyces lividans	Streptomyces lividans	Methylococcus capsulatus	Chloroplast Arabidopsis thaliana	Homo sapiens	Homo sapiens	Mycobacterium tuberculosis	Methylobacterium extorquens	Caulobacter crescentus	Streptomyces roseofulvus	Drosophila melanogaster	Drosophila melanogaster	Saccharopolyspora	Mycobacterium tuberculosis	Mycobacterium tuberculosis	Homo sapiens Homo sapiens	Homo sapiens
Table 4 (continued) S.lividans i protein and ATP synthase genes.	Mycobacterium tuberculosis H37Rv complete genome; segment 57/162.	Mycobacterium leprae cosmid L471.	S.lividans i protein and ATP synthase genes.	S.lividans i protein and ATP synthase genes.	M.capsulatus orfx, orfz, sqs and shc genes.	Arabidopsis thaliana chloroplast genomic DNA, complete sequence, strain:Columbia.	Homo sapiens clone RP11-114I16, *** SEQUENCING IN PROGRESS ***, 39 unordered pieces.	Homo sapiens clone RP11-114I16, *** SEQUENCING IN PROGRESS ***, 39 unordered pieces.	Mycobacterium tuberculosis H37Rv complete genome; segment 47/162.	Methylobacterium extorquens methanol oxidation genes, glmU-like gene, partial cds, and orf.2, orft.1, orfR genes, complete cds.	C.crescentus flagellar gene promoter region.	Streptomyces roseofulvus frenolicin biosynthetic gene cluster, complete sequence.	Drosophila melanogaster chromosome 2 clone BACR04B09 (D576) RPCI-98 04.B.9 map 43E1244F1 strain y; cn bw sp, *** SEQUENCING IN PROGRESS ***, 150 unordered pieces.	Drosophila melanogaster chromosome 2 clone BACR04B09 (D576) RPCI-98 04.B.9 map 43E12-44F1 strain y; cn bw sp. *** SEQUENCING IN PROGRESS ***. 150 unordered places.	Saccharopolyspora erythraea ferredoxin (fdxA) gene, complete cds.	Mycobacterium tuberculosis H37Rv complete genome; segment 51/162.	Mycobacterium tuberculosis sequence from clone y348.		unordered pieces. Homo sapiens clone NH0122L09, *** SEQUENCING IN PROGRESS ***, 2 unordered pieces.
222606	273419	U15186	222606	222606	Y09978	154478 AP000423	164070 AC009762	164070 AC009762	292539	AF017435	M69228	AF058302	165741 AC007301	165741 AC007301	M61119	AL010186	AD000020	174503 AC005697 160723 AC010722	160723 AC010722
8560	35516	36241	8560	8560	5538	154478	164070	164070	38970	4301	4424	25306	165741	165741	3869	37840	40056	174503 160723	160723
GB_BA1:SLATPSYNA	GB_BA1:MTCY373	GB_BA1:MLU15186	GB_BA1:SLATPSYNA	GB_BA1:SLATPSYNA	GB_BA1:MCSQSSHC	GB_PL1:AP000423	GB_HTG6:AC009762	GB_HTG6:AC009762	GB_BA1:MTCY10G2	GB_BA2:AF017435	GB_BA1:CCRFLBDBA	GB_BA2:AF058302	GB_HTG3:AC007301	GB_HTG3:AC007301	GB_BA1:SERFDXA	GB_BA1:MTV005	GB_BA1:MSGY348	GB_PR3:AC005697 GB_HTG3:AC010722	GB_HTG3:AC010722
rxa01201 1764			xa01202 1098			rxa01204 933			rxa01216 1124			rxa01225 1563	·		rxa01227 444			rxa01242 900	

WO 01/00844

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23-OCT-1998	23-Nov-98	23-Nov-98	1-Feb-99	16-OCT-1999	16-OCT-1999	29-OCT-1999			5-Nov-97	21-MAR-1997	18-Apr-98	25-Sep.00	25_Sen.99		23-Jun-99	01-MAR-1994	2-Nov-99	2-Nov-99	2-Nov-99	07-DEC-1999	07-DEC-1999
38,722	35,448	35,694	100,000	37,178	37,178	59.719	!		59,735	37,904	37,340	36 385	36.385	30 404	46,252	46,368	36,016	36,016	39,618	35,366	35,366
Magnaporthe grisea	Caenorhabditis elegans	Caenorhabditis elegans	Corynebacterium	Brosophila melanogaster	Drosophila melanogaster	r Escherichia coli			l) Escherichia coli	Escherichia coli	Homo sapiens	Homo sapiens	Homo sapiens	Fecharichia coli	Mycobacterium	tuberculosis Mycobacterium leprae	Homo sapiens	Homo sapiens	Homo sapiens	Drosophila melanogaster	Drosophila melanogaster
	Caenorhabditis elegans cosmid K05D4, complete sequence.	Caenorhabditis elegans cosmid K05D4, complete sequence.	confined accentum gradamicam pagene, complete CDS.				Wzx (wzx), WbnA (wbnA), O-antigen polymerase Wzy (wzy), WbnB (wbnB), WbnC (wbnC), WbnD (wbnD), WbnE (wbnE), UDP-Gic-4-epimerase GalE (galE), 6-phosphogluconate dehydrogenase Gnd (gnd), UDP-Gic-6.	dehydrogenase Ugd (ugd), and WonF (wbnF) genes, complete cds, and chain length determinant Wzz (wzz) gene, partial cds.	Escherichia coli hypothetical uridine-5'-diphosphoglucose dehydrogenase (ugd) Escherichia coli and O-chain length requiator (way) negas complete add	E.coli genomic DNA, Kohara clone #351(45.1-45.5 min.).			unordered pieces. Howa10K15, *** SEQUENCING IN PROGRESS ***, 4	unordered pieces. Escherichia coli K-12 MG1655 section 377 of 400 of the complete nenome	Mycobacterium tuberculosis H37Rv complete genome; segment 143/162.	Mycobacterium leprae cosmid L308.	Homo sapiens chromosome 7, *** SEQUENCING IN PROGRESS ***, 24 unordered nieces	Homo sapients chromosome 7, *** SEQUENCING IN PROGRESS ***, 24	Interest process of the second	Drosophila melanogaster chromosome 2 clone BACR03D06 (D569) RPCI-98 03.D.6 map 32A-32A strain y; cn bw sp, *** SEQUENCING IN PROGRESS***	91 unordered pieces. Drosophila melanogaster chromosome 2 clone BACR19N18 (D572) RPCI-98 19.N.18 map 32A-32A strain y; cn bw sp, *** SEQUENCING IN PROGRESS ***, 22 unordered pieces.
AQ255057	Z92804	292804 Y16642	7	AC010567	143287 AC010567	AF172324			U78086	D90841	144368 AC004103	215529 AC007383	AC007383	AE000487	AL021841	U00022	215767 AC009245	215767 AC009245	AC009245	225851 AC007186	202291 AC007147
583	19000	1800		143287	143287	14263			4759	20226	144368	215529	215529	13889	53662	36411	215767	215767	215767	225851	202291
GB_GSS10:AQ255057 583	GB_IN1:CEK05D4	GB_BA1:CGLPD		GB_HTG4:AC010567	GB_HTG4:AC010567	GB_BA2:AF172324			GB_BA2:ECU78086	GB_BA1:D90841	GB_PR3:AC004103	GB_HTG3:AC007383	GB_HTG3:AC007383	GB_BA2:AE000487	GB_BA1:MTV016	GB_BA1:U00022	GB_HTG4:AC009245	GB_HTG4:AC009245	GB_HTG4:AC009245	GB_HTG6:AC007186	GB_HTG6:AC007147
rxa01243 1083		rxa01259 981				xa01262 1284					rxa01311 870			rxa01312 2142			rxa01325 795			rxa01332 576	·

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16-Sep-99	•	19-OCT-1999	25-Feb-99	25-Feb-99	17-Jun-98		26-Apr-93		20-Apr-99	17-Jun-98		14-MAY-1997	14-MAY-1997		10-Feb-99	16-Nov-99		16-Nov-99		8JUL-8		24-Feb-97	00 00	90-100 100-100	23-MAR-1999		29-MAY-1997	29-Sep-97	9-Apr-97 3-Jun-99
34.821		58,487	37,963	37,963	38,011		47,726		36,599	36.940		35,284	38,324		39,778	32,658		38,395		122,66		100,000	26 766	000	100,000		53,041	54,461	39,286 39,412
5 Homo sapiens	-	Aquaspirillum arcticum	Caenorhabditis elegans	Caenorhabditis elegans	Mycobacterium	tuberculosis	Xanthomonas campestris		Homo sapiens	Mycobacterium	tuberculosis	Arabidopsis thaliana	Arabidopsis thaliana		Mycobacterium tuberculosis	Homo sapiens		Homo sapiens	:	Streptomyces lividans		Corynebacterium	glutamicum	como sapiens	Corynebacterium	glutamicum	Escherichia coli	Escherichia coli	Helicobacter pylori Homo sapiens
Table 4 (continued) Homo sapiens clone RPCI11-375120 SEQUENCING IN PROGRESS 25 Homo sapiens	unordered pieces.	Aquaspirillum arcticum malate dehydrogenase (MDH) gene, complete cds.	Caenorhabditis elegans clone Y40G12, *** SEQUENCING IN PROGRESS***, 8 unordered pieces.	Caenorhabditis elegans clone Y40G12, *** SEQUENCING IN PROGRESS***, 8 unordered pieces.	Mycobacterium tuberculosis H37Rv complete genome; segment 139/162.		Xanthomonas campestris phosphoglucomutase and phosphomannomutase	(xank) and phosphomannose isomerase and GDP-mannose pyrophosphorylase (xanB) genes, complete cds.	RPC111-47D24.TJ RPCI-11 Homo sapiens genomic clone RPCI-11-47D24,	genomic survey sequence. Mycobacterium tuberculosis H37Ry complete genome: segment 139/162.		T27A19-T7 TAMU Arabidopsis thaliana genomic clone T27A19, genomic	survey sequence. T21A19-T7.1 TAMU Arabidopsis thaliana genomic clone T21A19, genomic	survey sequence.	Mycobacterium tuberculosis H3/Rv complete genome; segment 141/162.	Homo sapiens clone RP11-252018, WORKING DRAFT SEQUENCE, 121	unordered pieces.	Homo sapiens clone RP11-252O18, WORKING DRAFT SEQUENCE, 121	unordered pieces.		transcriptional regulator, putative ferredoxin, putative cytochrome P450 oxidoreductase, and putative oxidoreductase genes, complete cds; and	C.glutamicum lysE and lysG genes.	Home conjust 19413 3 B&C BBC111 428A30 (Because Book Assessed	Institute Human BAC Library) complete sequence.	C.glutamicum pta gene and ackA gene.	•	E.coli genomic DNA, Kohara clone #405(52.0-52.3 min.).	DNA encoding acetate kinase protein form Escherichia coli.	Helicobacter pylori feoB-like DNA sequence, genomic survey sequence. we81c04.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone IMAGE:2347494 3' similar to gb:L19686_ma1 MACROPHAGE MIGRATION INHIBITORY FACTOR (HUMAN);, mRNA sequence.
207890 AC010207	1		AC006759	103725 AC006759	Z95121		M83231		AQ194038	295121	!	B10037	B09549		292771	262181 AC007547		262181 AC007547		AF0/2/09		X96471	185057 ACOUSOUS		X89084		D90861	E02087	U60627 AI701691
207890	;	066	103725	103725	36330	!	3410		269	36330	į	974	1097	1	42729	262181		262181	000	9300		2374	185052	70000	3657		14839	1200	349 349
GB HTG3:AC010207		GB_BA2:AF109682	GB_HTG2:AC006759	GB_HTG2:AC006759	GB_BA1:MTY20B11		GB_BA1:XANXANAB		GB_GSS10:AQ194038	GB_BA1:MTY20B11		GB_GSS3:B10037	GB_GSS3:B09549		GB_BA1:MICY/1	GB_HTG5:AC007547		GB_HTG5:AC007547	0010101010100	GB_BAZ:AFU/2/U9		GB_BA1:CGLYSEG	SP DD4.ACODEOUR	מפריים ביולים	GB BA1:CGPTAACKA	1	GB_BA1:D90861	GB_PAT:E02087	GB_GSS1:HPU60627 GB_EST31:Al701691
		rxa01350 1107			rxa01365 1497					rxa01369 1305					xa013// 1209				0007	Mau 1382 1200					xa01436 1314			:	ra01468 948

WO 01/00844

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14-Aug-97 16-Aug-99	05-MAY-1999	9-Apr-97	20-Aug-98 01-MAR-1994	18-Jun-98	27-Aug-99 24-Jun-99	23-OCT-1996	7-Feb-99 1-Jul-99	27-Apr-99		5-Feb-99	28-OCT-1998 15-OCT-1998	17-Jun-98	4-Nov-98
39,574	38,126	41,852	62,149 38,303	38,179	66,208 38,553	52,690	56,487 55,100	56,708		44,050	38,587 38,621	59,035	59,714
Homo sapiens Streptomyces coelicolor	A3(2) Streptomyces coelicolor	Corynebacterium	glutamicum Streptomyces coelicolor Mycobacterium leprae	Mycobacterium tuberculosis	Mycobacterium leprae Mycobacterium	tuberculosis Pseudomonas aeruginosa	Synechocystis sp. Drosophila melanogaster	s. Drosophila melanogaster		Streptococcus mutans	Streptomyces venezuelae Streptomyces venezuelae	Mycobacterium	tuberculosis Klebsiella pneumoniae
Table 4 (continued) ne31f04.s1 NCI_CGAP_Co3 Homo sapiens cDNA clone IMAGE:898975.3' similar to gb:L19686_rna1 MACROPHAGE MIGRATION INHIBITORY FACTOR (HUMAN);, mRNA sequence. Streptomyces coelicolor cosmid 151.	Streptomyces coelicolor cosmid E36.	Corynebacterium glutamicum multidrug resistance protein (cmr) gene,	Complete cas. Streptomycas coelicolor cosmid 6G4. Mycobacterium leprae cosmid B229.	Mycobacterium tuberculosis H37Rv complete genome; segment 146/162.	Mycobacterium leprae cosmid B1222. Mycobacterium tuberculosis H37Rv complete genome; segment 48/162.	Pseudomonas aeruginosa fumarase (fumC) and Mn superoxide dismutase	(sod.A) genes, complete cds. Synechocystis sp. PCC6803 complete genome, 9/27, 1056467-1188885. Drosophila melanogaster glycogen phosphorylase (GlyP) gene, complete cds.	Drosophila melanogaster glycogen phosphorylase (Glp1) mRNA, complete cds. Drosophila melanogaster		Streptococcus mutans DNA for dTDP-rhamnose synthesis pathway, complete	cas. Streptomyces venezuelae pikCD operon, complete sequence. Streptomyces venezuelae cytochrome P450 monooxygenase (pick) gene,	complete cos. Mycobacterium tuberculosis H37Rv complete genome; segment 16/162.	Klebsiella pneumoniae dTDP-D-glucose 4,6 dehydratase (rmlB), glucose-1-phosphate thymidylyl transferase (rmlA), dTDP-4-keto-L-rhamnose reductase (rmlD), dTDP-4-keto-6-deoxy-D-glucose 3,5-epimerase (rmlC), and rhamnosyl transferase (wbbL) genes, complete cds.
AA480256 AL109848	AL049763	U43535	AL031317 U00020	295389	AL049491 AL021897	U72494	D90907 AF073177	AF073179		D78182	AF079139 AF087022	Z96800	AF097519
389 40745	12581	2531	41055 36947	22255	34714 67200	4368	132419 9534	3159		7836	4342 1470	38900	4594
GB_EST15:AA480256 GB_BA1:SCI51	GB_BA1:SCE36	GB_BA1:CGU43535	GB_BA1:SC6G4 GB_BA1:U00020	GB_BA1:MTCY77	GB_BA1:MLCB1222 GB_BA1:MTV017	GB_BA1:PAU72494	GB_BA1:D90907 GB_IN2:AF073177	GB_IN2:AF073179		GB_BA1:D78182	GB_BA2:AF079139 GB_BA2:AF087022	GB_BA1:MTCY63	GB_BA2:AF097519
rxa01478 1959			rxa01482 1998	rxa01534	rxa01535 1530		rxa01550 1635		гха01562	xa01569 1482		rxa01570 978	

				Table 4 (continued)				
	GB_BA2:NGOCPSPS	8905	L09189	ts (rfbB), glucose-1- i, complete cds and UPD-	Neisseria meningitidis	58,384	30-Jul-96	*****
rxa01571 723	GB_BA1:AB011413	12070	AB011413	3, Orf4, Orf5, AfsA, Orf8, partial and	Streptomyces griseus 5	57,500	7-Aug-98	,000
	GB_BA1:AB011413	12070	AB011413	yces griseus genes for Orf2, Orf3, Orf4, Orf5, AfsA, Orf8, partial and cds.	Streptomyces griseus	35,655	7-Aug-98	**
rxa01572 615	GB_BA1:AB011413	12070	AB011413	Streptomyces griseus genes for Orf2, Orf3, Orf4, Orf5, AfsA, Orf8, partial and Streptomyc complete cds.	Streptomyces griseus 5	57,843	7-Aug-98	
	GB_BA1:AB011413	12070	AB011413	yces griseus genes for Orf2, Orf3, Orf4, Orf5, AfsA, Orf8, partial and cds.	Streptomyces griseus	38,119	7-Aug-98	
rxa01606 2799	GB_VI:CFU72240	4783	U72240	Choristoneura fumiferana nuclear polyhedrosis virus ETM protein homolog, 79 Choristone KDa protein homolog, 15 kDa protein homolog and GTA protein homolog nucleopoly	Choristoneura fumiferana 3 nucleopolyhedrovirus	37,115	29-Jan-99	
	GB_GSS10.AQ213248 408	408	AQ213248	Spring, 2011-202, MR CIT Approved Human Genomic Sperm Library D Homo Homo sapiens sapiens genomic clone Plate=3249 Col=3 Row=B, genomic survey sequence.		34,559	18-Sep-98	
	GB_GSS8:AQ070145	285	AQ070145	HS_3027_B1_H02_MR CIT Approved Human Genomic Sperm Library D Homo Homo sapiens sapiens genomic clone Plate=3027 Col=3 Row=P, genomic survey sequence.		40,351	5-Aug-98	104
rxa01626 468	GB_PR4:AF152510	2490	AF152510	Homo sapiens protocadherin gamma A3 short form protein (PCDH-gamma-A3) Homo sapiens variable region sequence complete cds		34,298	14-Jul-99	
	GB_PR4:AF152323	4605	AF152323	Homo sapiens protocadherin gamma A3 (PCDH-gamma-A3) mRNA, complete Homo sapiens		34,298	22-Jul-99	
ma01632 1128	GB_PR4:AF152509 GB_HTG4:AC006590	2712 127171	AF152509 AC006590	o sapiens PCDH-gamma-A3 gene, aberrantly spliced, mRNA sequence. ophila melanogaster chromosome 2 clone BACR13N02 (D543) RPCI-98 map 36E-36E strain y; cn bw sp, *** SEQUENCING IN PROGRESS***, innerdend nieces.	lanogaster	34,298 33,812	14-Jul-99 19-OCT-1999	
	GB_HTG4:AC006590	127171	127171 AC006590	ter chromosome 2 clone BACR13N02 (D543) RPCI-98 strain y; cn bw sp, *** SEQUENCING IN PROGRESS***,	Drosophila melanogaster	33,812	19-OCT-1999	
	GB_GSS8:B99182	415	B99182	CIT-HSP-2280113.TX CIT-HSP Homo sapiens genomic clone 2280113, Homo sapiens panomic survey sequence		36,111	26-Jun-98	
rxa01633 1206	GB_BA1:BSUB0009	208780	Z99112	Bacillus subtilis complete genome (section 9 of 21): from 1598421 to 1807200. Bacillus subtilis		36,591	26-Nov-97	.,
	GB_BA1:BSUB0009	208780	299112	Bacillus subtilis complete genome (section 9 of 21): from 1598421 to 1807200. Bacillus subtilis		34,941	26-Nov-97	
	GB_HTG2:AC006247	174368	174368 AC006247	Drosophila melanogaster chromosome 2 clone BACR48110 (D505) RPCI-98 Drosophila 48.I.10 map 49E6-49F8 strain y; cn bw sp, *** SEQUENCING IN PROGRESS ***, 17 unordered pieces.	Drosophila melanogaster	37,037	2-Aug-99	

WO 01/00844

	11-Aug-98	17-Jun-98	18-Jul-95	12-Sep-93	17-Jun-98	27-Aug-99	13-Jul-95	03-DEC-1998	27-Aug-98	17-Jun-98	31-Jul-97		26-Nov-98	. 86-unr-/L	22-Aug-97	12-Nov-98	23-Nov-99	5-Sep-97	29-Aug-97	21-Sep-94	17-Jun-98	17-Jun-98	24-Sep-98	30-OCT-1998	7-Feb-99
	100,000	38,626	36,783	99,913	38,786	38,238	35,334	39,222	40,653	36,650	63,438	000	53,088 53,088	£2,081	61,364	52,323	39,209	9 40,021	e 34,375	62,173	39,749	40,034	38,068	36,557	35,316
	Corynebacterium	glutamicum Mycobacterium tuberculosis			Mycobacterium tuberculosis	Mycobacterium leprae	Caenorhabditis elegans	Populus balsamifera subso trichocama		Mycobacterium tuberculosis	Mycobacterium	Tuberculosis	Streptomyces coelicolor	Mycobacterium tuberculosis	Mycobacterium leprae	Escherichia coli	Homo sapiens	Saccharomyces cerevisiae 40,021	Saccharomyces cerevisiae 34,375	Acetobacter xylinus	Mycobacterium tuberculosis	Mycobacterium tuberculosis	Homo sapiens	Drosophila melanogaster	'Homo sapiens
Table 4 (continued)	Corynebacterium glutamicum DNA for L-Malate:quinone oxidoreductase.	Mycobacterium tuberculosis H37Rv complete genome; segment 124/162.	Drosophila melanogaster kinesin-like protein (klp68d) mRNA, complete cds.	Corynebacterium glutamicum fda gene for fructose-bisphosphate aldolase (EC 4.1.2.13).	Mycobacterium tuberculosis H37Rv complete genome; segment 18/162.	Mycobacterium leprae cosmid B4.	Caenorhabditis elegans cosmid C27H5.	xylem.est.878 Poplar xylem Lambda ZAPII library Populus balsamifera subsp. trichocarpa cDNA 5', mRNA sequence.		Mycobacterium tuberculosis H37Rv complete genome; segment 72/162.	Mycobacterium tuberculosis cytochrome D oxidase subunit I (appC) gene,	Strentomyces coelicator coemid D28	Outprofit year openical of the Dro. Mycobacterium tubercuiosis H37Dv complete pename: somment 081162	my condecendin tabel canonis i 137 NV complete genome, segment 96/102.	Mycobacterium leprae cosmid B22.	Escherichia coli K-12 MG1655 section 65 of 400 of the complete genome.	Human DNA sequence from BAC 57G9 on chromosome 22q12.1 Contains ESTs, CA repeat, GSS.	Saccharomyces cerevisiae chromosome VIII cosmid 9666.	Saccharomyces cerevisiae chromosome VIII cosmid 9986.	Acetobacter xylinum phosphoglucomutase (celB) gene, complete cds.	Mycobacterium tuberculosis H37Rv complete genome; segment 133/162.	Mycobacterium tuberculosis H37Rv complete genome; segment 133/162.	HS_2222_B1_H03_MR CIT Approved Human Genomic Sperm Library D Homo Homo sapiens sapiens genomic clone Plate=2222 Col=5 Row=P, genomic survey sequence.	Drosophila melanogaster, chromosome 2L, region 30A3- 30A6, P1 clones DS06958 and DS03097, complete sequence	Homo sapiens genomic DNA, 21q region, clone: B137B7BB68, genomic survey Homo sapiens sequence.
	AJ224946	295207	U15974	X17313	Z95324	AL023514	U14635	Al167112	AQ102635	Z95554	AF009226	AI 034355	770283		Z98741		295116	U10397	U00027	L24077	Z83866	Z83866	AQ142579	108924 AC005889	AG008814
	2408	20270	2994	3371	35019	36310	35840	579	347	35938	965	36224	34150		40281	15067	1138/2	39057	41664	2058	31859	31859	529	108924	637
	GB_BA1:CGA224946	GB_BA1:MTCY24A1	GB_IN1:DMU15974	GB_BA1:CGFDA	GB_BA1:MTY13E10	GB_BA1:MLCB4	GB_INZ:CELC27H5	GB_ES124:Al167112	GB_GSS9:AQ102535	GB_BA1:MTCY01B2	GB_GSS1:AF009226	GB BA1:SCD78	GB_BA1:MTCY190		GB_BA1:MLCB22	GB_BAZ:AE000175	GB_PK3:H55/G9	GB_PL2:YSCH9666	GB_PL2:YSCH9986	GB_BA1:ABCCELB	GB_BA1:MTCY22D7	GB_BA1:MTCY22D7	GB_GSS9:AQ142579	GB_IN2:AC005889	GB_GSS1:AG008814
	rxa01695 1623			CaU1/02 1155			rxa01743 901			rxa01744 1662			rxa01745 836			01160 4140				rxa01814 1785			rxa01851 1809		

WO 01/00	844						106							PCT	/ IB 00/009	43
03-OCT-1999	15-Nov-99	03-OCT-1999	13-MAR-1996	17-Jun-98	10-DEC-1996	27-Apr-93 23-Nov-98	18-OCT-1997	17-Jun-98	29-MAR-1999	7-Feb-99 20-Apr-99	00 411	28-Nov-98	03-DEC-1999	03-DEC-1999	01-OCT-1999	2-Aug-99
36,364	35,334	36,529	59,862	61,949	806'69	36,899 36,899	34,805	37,892	40,413	47,792	30 306	42,807	36,417	37,667	39,640	32,969
Microcystis aeruginosa	Trypanosoma brucei	Microcystis aeruginosa	Saccharopolyspora erythraea	Mycobacterium tuberculosis	Mycobacterium tuberculosis	Homo sapiens Homo sapiens	Homo sapiens	Mycobacterium tuberculosis	Streptomyces coelicolor	Synechocystis sp. Homo sapiens	Themotoment	Drosophila melanogaster	Homo sapiens	Homo sapiens	Acinetobacter Iwoffii	Drosophila melanogaster
Microcystis aeruginosa DNA polymerase III beta subunit (dnaN) gene, partial cds; microcystin synthetase gene cluster, complete sequence; Uma1 (uma1), Uma2 (uma2), Uma3 (uma3), Uma4 (uma4), and Uma5 (uma5) genes, complete cds; and Uma6 (uma6) gene, partial cds.	Trypanosoma brucei chromosome II clone RPCI93-25N14, *** SEQUENCING IN PROGRESS ***, 2 unordered pieces.	Microcystis aeruginosa DNA polymerase III beta subunit (dnaN) gene, partial cds; microcystin synthetase gene cluster, complete sequence; Uma1 (uma1), Uma2 (uma2), Uma3 (uma3), Uma4 (uma4), and Uma5 (uma5) genes, complete cds; and Uma6 (uma6) gene, partial cds.	Saccharopolyspora erythraea ferredoxin (fdxA) gene, complete cds.	Mycobacterium tuberculosis H37Rv complete genome; segment 51/162.	Mycobacterium tuberculosis sequence from clone y348.	Human kidney alpha-2-adrenergic receptor mRNA, complete cds. Homo sapiens alpha2-C4-adrenergic receptor gene, complete cds.	HS-1055-B1-A03-MR.abi CIT Human Genomic Sperm Library C Homo sapiens Homo sapiens genomic clone Plate=CT 777 Col=5 Row=B, genomic survey sequence.	Mycobacterium tuberculosis H37Rv complete genome; segment 69/162.	Streptomyces coelicolor A3(2), glycogen metabolism cluster II.	Synechocystis sp. PCC6803 complete genome, 10/27, 1188886-1311234. RPCI11-49P6.TK.1 RPCI-11 Homo sapiens genomic clone RPCI-11-49P6,	genomic survey sequence. Thermotona maritima section 33 of 136 of the complete genome	GM01044.5prime GM Drosophila melanogaster ovary BlueScript Drosophila melanogaster con CDNA clone GM01044 5prime. mRNA sequence.	Homo sapiens clone RP3-405J10, *** SEQUENCING IN PROGRESS ***, 102 unordered places.	Homo sapiens clone RP3-405J10, *** SEQUENCING IN PROGRESS ***, 102 unordered pieces.	Acinetobacter lwoffii wzc, wzb, wza, weeA, weeB, wceC, wzx, wzy, weeD, weeE, weeG, weeH, weeI, weeV, weeK, galU, ugd, pgi, galE, pgm (partial) and mip (partial) genes (emulsan biosynthetic gene cluster), strain RAG-1.	Drosophila melanogaster chromosome 3 clone BACR02L12 (D753) RPCI-98 02.L.12 map 94B-94C strain y; cn bw sp, *** SEQUENCING IN PROGRESS*** 113 unordered pieces.
AF183408	158889 AC008031	AF183408	M61119	AL010186	AD000020	J03853 U72648	B42200	Z74020	AJ001206	D90908 AQ116291	AE001721	AA567090	303147 AC008147	303147 AC008147	AJ243431	125235 AC008197
63626	_	. 0	3869	37840	40056	1491 4850	387	35377	9184	122349 D90908 572 AQ1162	17632	296	303147	303147	26953	125235
GB_BA2:AF183408	GB_HTG5:AC008031	GB_BA2:AF183408	GB_BA1:SERFDXA	GB_BA1:MTV005	GB_BA1:MSGY348	GB_PR1:HUMADRA2C 1491 GB_PR4:HSU72648 4850	GB_GSS3:B42200	GB_BA1:MTCY48	GB_BA1:SCO001206	GB_BA1:D90908 GB_GSS9:AQ116291	GB BA2:AE001721	GB_EST16:AA567090	GB_HTG6:AC008147	GB_HTG6:AC008147	GB_BA2:ALW243431	GB_HTG2:AC008197
rxa01859 1050			rxa01865 438			rxa01882 1113		rxa01884 1913	-	rxa01886 897			rxa01887 1134			rxa01888 658

WO 01/00844								107								PCT/IB00/00943									
2-Aug-99	21-Jul-99	05-MAR-1999	1-Aor-97	2-Jun-98	24-Apr-99	05-MAY-1998	17-Jun-98	6-Sep-94	17-MAR-1999	02-OCT-1998	11-MAY-1999	04-MAY-1992	24-Aug-99	25-MAR-1998	16-MAR-1999	25-Nov-98	30 Apr 06	06-1d-07	26-Nov-97	28-Sep-99	28-Nov-96	26-Nov-97	2-Nov-93	17-Jun-98	
32,969	43,617	40,040	37.844	37,136	100,000	s 65,254	40,058	59,551	39,468	39,291	38,384	56,283	37,593	36,309	41,941	39,855	66 202	707,00	37,255	63,607	877 78	35,574	51.826	54,476	
Drosophila melanogaster	Zea mays	Human immunodeficiency	virus type 1 Aspergillus fumigatus	Homo sapiens	Corynebacterium	glutamicum Mycobacterlum smegmatis 65,254	Mycobacterium	tuberculosis Mycobacterium leprae	Streptomyces coelicolor	Homo sapiens	Corynebacterium glutamicum	. Arthrobacter sp.	Homo sapiens	Aquifex aeolicus	Drosophila melanogaster	Drosophila melanogaster	Bacillus	stearothermophilus	Bacillus subtilis	Streptococcus mutans	Staphylococcus xylosus	Bacillus subtilis	Bacillus subtilis	Mycobacterium tuberculosis	
Table 4 (continued) Drosophila melanogaster chromosome 3 clone BACR02L12 (D753) RPCI-98 02.L.12 map 948-94C strain y; cn bw sp. *** SEQUENCING IN PROGRESS ***, 113 unordered pieces.	606070C09.y1 606 - Ear tissue cDNA library from Schmidt lab Zea mays cDNA, Zea mays	minory sequence. Human immunodeficiency virus type 1 subtype C nef gene, patient MP83.	A.fumigatus chsE gene.	Homo sapiens chromosome 21q22.3 BAC 28F9, complete sequence.	Corynebacterium glutamicum non gene.	Mycobacterium smegmatis NADH dehydrogenase (ndh) gene, complete cds.	Mycobacterium tuberculosis H37Rv complete genome; segment 84/162.	M. leprae genomic DNA sequence, cosmid B38 bfr gene, complete cds.		Homo sapiens chromosome 7qtelo BAC E3, complete sequence. Coluiamicum nanB nanC & volB canae	Colored Period Caylor general	Arthrobacter Sp. N.R.R.L. B3728 xylA gene for D-xylose(D-glucose) isomerase. Arthrobacter sp.	Homo sapiens clone NH0511A20, *** SEQUENCING IN PROGRESS ***, 6 unordered pieces.	Aquifex agolicus section 71 of 109 of the complete genome.	LD39282.5prime LD Drosophila melanogaster embryo pOT2 Drosophila melanonaster cDNA close I D19282 Eccino. mDNA consessor	LD2827. Spring LD Drosophila melanogaster enbryo pOT2 Drosophila melanogaster enbryo pOT2 Drosophila	Bacillus stearothermophilus paid gene for phosphoplucoisomerase isoenzyme	A (EC 5.3.1.9).	Bacillus subtilis complete genome (section 17 of 21); from 3197001 to 3414420.	Streptococcus mutans sorbitol phosphoenolpyruvate:sugar phosphotransferase Streptococcus mutans operon: complete sequence and unknown gene	S.xylosus scrB and scrR genes.	Bacillus subtilis complete genome (section 20 of 21): from 3798401 to	40 tubbu. B.subtilis genomic region (325 to 333).	Mycobacterium tuberculosis H37Rv complete genome; segment 46/162.	
125235 AC008197	AI881527	AJ232971	Y09542	AF064858	A3230230	AF038423	Z83859	L01095 Al 035640	20000	AF093117 X96580		X59466	176060 AC009500	AE000739	AI519629	AA949396	X16639		Z99120	AF132127	X67744	Z99123	X73124	Z 94752	
125235	298	621	6158	193387	280	1376	36021	37114	3	147216 2164		1905	176060	13335	612	167	1822		217420	8452	3161	212150	97015	27030	
GB_HTG2:AC008197	GB_EST36:AI881527	GB_VI:HIV232971	GB_PL1:AFCHSE	GB_PR3:AF064858	00700700700	GB_BA2:AF038423	GB_BA1:MTCY359	GB_BA1:MSGB38COS GB_BA1:SCE63		GB_PR3:AF093117 GB_BA1:CGPAN		GB_BA1:ASXYLA	GB_HTG3:AC009500	GB_BA2:AE000739	GB_EST28:AI519629	GB_EST21:AA949396	GB_BA1:BSPGIA		GB_BA1:BSUB0017	GB_BA2:AF132127	GB_BA1:SXSCRBA	GB_BA1:BSUB0020	GB_BA1:BSGENR	GB_BA1:MTCl237	
		rxa01891 887		CX301895 1051	200			xa01901 1383		xa01927 1503				rxa01952 1836			rxa01989 630				rxa02026 720			rxa02028 526	

1-Aug-97		29-Apr-99	27-Aug-99	06-107-11	14-Aug-97	28-Jul-99	6-Feb-99	; ;	18-Jun-98	08-DEC-1999	19-DEC-1997	04-MAR-1998		12-Jul-99	1-Jul-99	27-OCT-1999		10-DEC-1996	17-Jun-98		29-MAR-1999	29-MAY-1997	6-Sep-99	15-Sep-99	15-Jun-96	15-Jun-96	01-OCT-1999	17-Jun-98
36.100		9 32,039	61,896	† 6 6 6	59,659	98,928	98.928		39,265	37,453	37,711	37 711	5	56,972	40,696	36,795		40,156	55,218		38,475	38,586	37,259	38,868	51,399	51,399	36,683	57,292
Saccharomycas cerevisiae 36 100		, saccnaromyces cerevisia	Mycobacterium leprae	tuberculosis	Mycobacterium	Corynebacterium	glutamicum S Corynebacterium	glutamicum	Mycobacterium tuberculosis	Homo sapiens	Arabidopsis thaliana	Arabidopsis thaliana		Streptomyces coelicolor	Oryza sativa	Homo sapiens		Mycobacterium	tuberculosis Mycobacterium	tuberculosis	Streptomyces coelicolor	Escherichia coli	, Homo sapiens	Homo sapiens	Mycobacterium leprae	Mycobacterium leprae	Mus musculus	Mycobacterium tuberculosis
Table 4 (continued) Saccharomyces cerevisiae chromosome V cosmids 9537, 9581, 9495, 9867.	and lambda clone 5898.	V2009 III III-SXTAVIACZ, IIISETION LIDIAIY SACCHAIOMYCES CEFEVISIAE GENOMIC 5 , SACCHAIOMYCES CEFEVISIAE 3Z,U39 genomic survey sequence.	Mycobacterium leprae cosmid B1222.	my conduction is described and the complete gallonia, addition 1477 104.	Mycobacterium tuberculosis rfbA, rhamnose biosynthesis protein (rfbA), and	mino geries, complete cos. Brevibacterium lactofermentum gene for alpha-ketoglutaric acid	dehydrogenase. Corynebacterium glutamicum DNA for 2-oxoglutarate dehydrogenase, complete Corynebacterium	cds.	Mycobactenum tuberculosis H3/RV complete genome; segment 34/152.	Homo sapiens chromosome 17 clone RP11-958E11 map 17, ***	SEQUENCING IN PROGRESS ***, 2 ordered pieces. Arabidopsis thaliana chromosome II BAC T21L14 genomic sequence, complete Arabidopsis thaliana	sequence. Arabidoosis thaliana chromosome II BAC F2518 cenomic sequence, complete. Arabidoosis thaliana	sequence.	S.coelicolor DNA for glgC gene.	nbxb0074H11r CUGI Rice BAC Library Oryza sativa genomic clone	_	similar to WP:T03G11.6 CE04874;, mRNA sequence.	Mycobacterium tuberculosis sequence from clone y151.	Mycobacterium tuberculosis H37Rv complete genome; segment 59/162.		Streptomyces coelicolor A3(2) glycogen metabolism clusterl.	E.coli genomic DNA, Kohara clone #401(51.3-51.6 min.).	wq07d12.x1 NCI_CGAP_Kid12 Homo sapiens cDNA clone IMAGE:2470583 3', Homo sapiens mRNA sequence.	Homo sapiens chromosome 5 clone CITB-H1_2074D8, *** SEQUENCING IN PROGRESS ***, 77 unordered pieces.	Mycobacterium leprae cosmid B1551 DNA sequence.	Mycobacterium leprae cosmid B1554 DNA sequence.	Mus musculus transcription factor TBLYM (Tblym) mRNA, complete cds.	Mycobacterium tuberculosis H37Rv complete genome; segment 98/162.
018778	ACE04177	7 1000	AL049491 795390		U43540	E14601	D84102	900400	ALUZ 1000	211682 AC005883	AC003033	AC002334		X89733	AQ687350	AW028530		AD000018	Z73902		AJ001205	D90858	AI948595	220665 AC010387	L78813	L78814	AF093099	270283
66030			34714	2	3453	4394	4394	977	75440	211682	84254	75050		1518	786	444		37036	32514	;	9589	13548	469	220665	S 36548	S 36548	2482	34150
GB_PL2:SCE9537	T C C C C C C C C C C C C C C C C C C C	711000000000000000000000000000000000000	GB_BA1:MLCB1222		GB_BA1:MTU43540	GB_PAT:E14601	GB_BA1:D84102	900 EM-144	900A I MI 1 VO 20	GB_HTG7:AC005883	GB_PL2:ATAC003033	GB PL2:ATAC002334		GB_BA1:SCGLGC	GB_GSS4:AQ687350	GB_EST38:AW028530		GB_BA1:MSGY151	GB_BA1:MTCY130		GB_BA1:SCO001205	GB_BA1:D90858	GB_EST37:Al948595	GB_HTG3:AC010387	GB_BA1:MSGB1551CS 36548	GB_BA1:MSGB1554CS 36548	GB_RO:AF093099	GB_BA1:MTCY190
			rxa02054 1140			xa02056 2891				xa02061 1617				rxa02063 1350				rxa02100 2348				xa02122 822			rxa02140 1200			xa02142 774

WO 01/00844

	GB_BA1:SC6G10	36734	AL049497	Table 4 (continued) Streptomyces coelicolor cosmid 6G10.	Streptomyces coelicolor	35,058	24-MAR-1999
	GB_BA1:AB016787	5550	AB016787	Pseudomonas putida genes for cytochrome o ubiquinol oxidase A-E and 2	Pseudomonas putida	47,403	5-Aug-99
rxa02143 1011	GB_BA1:MTCY190	34150	270283	UKFS, complete cds. Mycobacterium tuberculosis H37Rv complete genome; segment 98/162.	Mycobacterium tuberculosis	57,317	17-Jun-98
	GB_BA1:MSGB1551CS 36548	S 36548	L78813	Mycobacterium leprae cosmid B1551 DNA sequence.	Mycobacterium leprae	38,159	15-Jun-96
785 AA120cm	GB_BA1:MSGB1554CS 36548	34460	770283	Mycobacterium leprae cosmid B1554 DNA sequence.	Mycobacterium leprae	38,159	15-Jun-96
1401 FF 208VI		<u> </u>	2 10283	Mycobacterium tuberculosis H3/RV complete genome; segment 98/162.	Mycobacterium tuberculosis	55,530	17-Jun-98
	GB_HTG3:AC011500_0 300851 AC011500	0 300851	AC011500	Homo sapiens chromosome 19 clone CIT978SKB_60E11, *** SEQUENCING IN PROGRESS ***, 246 unordered pieces.	Homo sapiens	39,659	18-Feb-00
	GB_HTG3:AC011500_0 300851 AC011500	0 300851	AC011500	Homo sapiens chromosome 19 clone CIT978SKB_60E11, *** SEQUENCING IN PROGRESS ***, 246 unordered pieces.	Homo sapiens	39,659	18-Feb-00
rxa02147 1140	GB_EST28:Al492095	485	AI492095	tg07a01.x1 NCI_CGAP_CLL1 Homo sapiens cDNA clone IMAGE:2108040 3', mRNA sequence.	Homo sapiens	39,798	30-MAR-1999
	GB_EST10:AA157467	376	AA157467	zo50e01.r1 Stratagene endothelial cell 937223 Homo sapiens cDNA clone	Homo sapiens	36,436	11-DEC-1996
	GB_EST10:AA157467	376	AA157467	zo50e01.r1 Stratagene endothelial cell 937223 Homo sapiens cDNA clone IMAGE:590328 5. mRNA sequence	Homo sapiens	36,436	11-DEC-1996
rxa02149 1092	GB_PR3:HSBK277P6	61698	AL117347	Human DNA sequence from clone 277P6 on chromosome 1q25.3-31.2,	Homo sapiens	36,872	23-Nov-99
	GB_BA2:EMB065R075		AF116423	complete sequence. Rhizobium etli mutant MB045 RosR-transcriptionally regulated sequence.	Rhizobium etli	43,175	06-DEC-1999
	GB_EST34:AI789323	574	AI789323	uk53g05.y1 Sugano mouse kidney mkia Mus musculus cDNA clone IMAGE:1972760 5' simitar to WP:K11H12.8 CE12160 ;, mRNA sequence.	Mus musculus	39,715	2-Jul-99
rxa02175 1416	GB_BA1:CGGLTG	3013	X66112	C.glutamicum glt gene for citrate synthase and ORF.	Corynebacterium	100,000	17-Feb-95
	- GB_BA1:MTCY31	37630	Z73101	Mycobacterium tuberculosis H37Rv complete genome; segment 41/162.	gradamicani Mycobacterium tuberculosis	64,331	17-Jun-98
CV307108 818	GB_BA1:MLCB57	38029	Z99494 M76426	Mycobacterium leprae cosmid 857.	Mycobacterium leprae	62,491	10-Feb-99
		2		complete cds.	Rattus norvegicus	18/,85	31-MAY-1995
	GB_GSS8:AQ012162	763	AQ012162	127PB037070197 Cosmid library of chromosome II Rhodobacter sphaeroides genomic clone 127PB037070197, genomic survey sequence.	Rhodobacter sphaeroides	40,044	4-Jun-98
	GB_RO:RATDAPRP	2819	M76426	Rattus norvegicus dipeptidyl aminopeptidase-related protein (dpp6) mRNA, complete cds.	Rattus norvegicus	37,312	31-MAY-1995
rxa02209 1694	GB_BA1:AB025424	2995	AB025424	Corynebacterium glutamicum gene for aconitase, partial cds.	Corynebacterium glutamicum	99,173	3-Apr-99
	GB_BA2:AF002133	15437	AF002133	Mycobacterium avium strain GIR10 transcriptional regulator (mav81) gene, partial cds, aconitase (acn), invasin 1 (inv1), invasin 2 (inv2), transcriptional regulator (moxR), ketoacyl-reductase (fabG), enoyl-reductase (inhA) and ferrochelatase (mav272) genes, complete cds.	Mycobacterium avium	40,219	26-MAR-1998

				Table 4 (continued)			
	GB_BA1:MTV007	32806	AL021184	AL021184 Mycobacterium tuberculosis H37Rv complete genome; segment 64/162.	Mycobacterium tuberculosis	38,253	17-Jun-98
rxa02213 874	GB_BA1:AB025424	2995	AB025424	Corynebacterium glutamicum gene for aconitase, partial cds.	Corynebacterium glutamicum	960'66	3-Apr-99
	GB_BA1:MTV007	32806	AL021184	Mycobacterium tuberculosis H37Rv complete genome; segment 64/162.	Mycobacterium tuberculosis	34,937	17-Jun-98
	GB_BA2:AF002133	15437	AF002133	Mycobacterium avium strain GIR10 transcriptional regulator (mav81) gene, partial cds, aconitase (acn), invasin 1 (inv1), invasin 2 (inv2), transcriptional regulator (moxR), ketoacyl-reductase (fabG), enoyl-reductase (inhA) and ferrochelatase (mav272) genes, complete cds.	Mycobacterium avium	36,885	26-MAR-1998
ка02245 780	GB_BA2:RCU23145	5960	U23145	Rhodobacter capsulatus Calvin cycle carbon dioxide fixation operon: fructose-1,5-/sedoheptulose-1,7-bisphosphate aldolase (cbbA) gene, partial cds, Form II ribulose-1,5-bisphosphate carboxylase/oxygenase (cbbM) gene, complete cds, and Calvin cycle operon: pentose-5-phosphate-3-epimerase (cbbE), phosphoglycolate phosphatase (cbbZ), and cbbY genes, complete cds.	Rhodobacter capsulatus	48,701	28-OCT-1997
	GB_BA1:ECU82664	139818	139818 U82664	Escherichia coli minutes 9 to 11 genomic sequence.	Escherichia coli	39,119	11-Jan-97
	GB_HTG2:AC007922	158858	158858 AC007922	Homo sapiens chromosome 18 clone hRPK.178_F_10 map 18, *** SEQUENCING IN PROGRESS ***, 11 unordered pieces.	Homo sapiens	33,118	26-Jun-99
rxa02256 1125	GB_BA1:CGGAPPGK	3804	X59403	C.glutamicum gap, pgk and tpi genes for glyceraldehyde-3-phosphate, ohospholycerate kinase and triosephosphate isomerase	Corynebacterium clutamicum	99,289	05-OCT-1992
	GB_BA1:SCC54	30753	AL035591	Streptomyces coelicolor cosmid C54.	Streptomyces coelicolor	36,951	11-Jun-99
	GB_BA1:MTCY493	40790	Z 95844	Mycobacterium tuberculosis H37Rv complete genome; segment 63/162.	Mycobacterium tuberculosis	64,196	19-Jun-98
rxa02257 1338	GB_BA1:CGGAPPGK	3804	X59403	C.glutamicum gap, pgk and tpi genes for glyceraldehyde-3-phosphate, phospholycerate kinase and triosephosphate isomerase.	Corynebacterium	98,873	05-OCT-1992
	GB_BA1:MTCY493	40790	295844	Mycobacterium tuberculosis H37Rv complete genome; segment 63/162.	Mycobacterium tuberculosis	61,273	19-Jun-98
	GB_BA2:MAU82749	2530	U82749	Mycobacterium avium glyceraldehyde-3-phosphate dehydrogenase homolog	Mycobacterium avium	61,772	6-Jan-98
rxa02258 900	GB_BA1:CGGAPPGK	3804	X59403	C.glutamicum gap, pgk and tpi genes for glyceraldehyde-3-phosphate,	Corynebacterium .	29,667	05-OCT-1992
				phosphoglycerate kinase and triosephosphate isomerase.	glutamicum		
	GB_BATICORPEPC	4885	818C2M	C.giutamicum pnospnoenoipyruvate carboxylase gene, complete cas.	Corynebacterium alutamicum	000,001	15-DEC-1995
	GB_PAT:A09073	4885	A09073	C.glutamicum ppg gene for phosphoenol pyruvate carboxylase.	Corynebacterium	100,000	25-Aug-93
rxa02259 2895	GB_BA1:CORPEPC	4885	M25819	C.glutamicum phosphoenolpyruvate carboxylase gene, complete cds.	glutamicum Corynebacterium	100,000	15-DEC-1995
	GB PAT:A09073	4885	A09073	C.clutamicum ppg gene for phosphoenol pyruvate carboxylase.	glutamicum Corynebacterium	100.000	25-Aug-93
	•				glutamicum		
	GB_BA1:CGPPC	3292	X14234	Corynebacterium glutamicum phosphoenolpyruvate carboxylase gene (EC 4.1.1.31).	Corynebacterium glutamicum	99,827	12-Sep-93

WO 01/00844

ка02404 2340 ка02414 870	GB_HTG2:AC007889 GB_BA1:CGACEA GB_BA1:CORACEA GB_BA1:CGACEB GB_BA1:CGACEB GB_BA1:CGACEB GB_BA1:CGACEB GB_BA1:CGACEB GB_BA1:CGACEB GB_BA1:CAC011214 GB_HTG3:AC011214		127840 AC007889 2427 X75504 1905 L28760 2135 I13693 3024 X78491 2725 L27123 5588 Y11998 176258 AC007102 183414 AC011214 7457 AF101055 4441 J03247 4458 M84656	Table 4 (continued) 127840 AC007889 Drosophila melanogaster chromosome 3 clone BACR48E12 (D695) RPCI-98 48.E.12 map 87A-87B strain y; cn bw sp, *** SEQUENCING IN PROGRESS***, 86 unordered pieces. 2427 X75504 C.glutamicum aceA gene and thiX genes (partial). 1905 L28760 Corynebacterium glutamicum isocitrate lyase (aceA) gene. 2135 113693 Sequence 3 from patent US 5438822. 2225 L27123 Corynebacterium glutamicum malate synthase (aceB) gene, complete cds. 2225 L27123 Corynebacterium glutamicum malate synthase (aceB) gene, complete cds. 2225 L27123 Corynebacterium glutamicum malate synthase (aceB) gene, complete cds. 2225 L27123 Corynebacterium glutamicum malate synthase (aceB) gene, complete cds. 2225 L27123 Corynebacterium glutamicum malate synthase (aceB) gene, complete cds. 2225 L27123 Corynebacterium glutamicum malate synthase (aceB) gene, complete cds. 2225 L27123 Corynebacterium glutamicum malate synthase (aceB) gene, complete cds. 2225 L27123 Corynebacterium glutamicum malate synthase (aceB) gene, complete sequence. Homo sapiens reading frames. 2326 L27123 Corynebacterium glutamicum adp operon, complete sequence. 2327 L27123 Corynebacterium glutamicum adp operon, complete sequence. 2328 Y11998 P.fluorescens FC2.1, FC2.2, FC2.3c, FC2.4 and FC2.5c open reading frames. 2329 Pseudomonas reading frames. 2441 J03247 AF101055 Clostridium acetobutylicum adp operon, complete sequence. 2441 J03247 Rabbit phosphorylase kinase (alpha subunit) mRNA, complete cds. 2448 M64656 Oryctolagus curiculus phosphorylase kinase alpha subunit mRNA, complete	Drosophila melanogaster 34,897 Corynebacterium 100,000 glutamicum 100,000 glutamicum 99,795 Gorynebacterium 99,795 glutamicum 99,786 glutamicum 99,786 glutamicum 99,786 Homo sapiens 35,069 Homo sapiens 35,885 Homo sapiens 36,885 Clostridium acetobutylicum 39,005 Oryctolagus cuniculus 36,000	34,897 100,000 100,000 99,795 99,786 63,539 63,539 35,069 36,885 36,885 36,061	2-Aug-99 9-Sep-94 10-Feb-95 13-Jan-95 11-Jul-97 2-Jun-99 03-OCT-1999 03-MAR-1999 22-Jun-98
ма02440 963 ма02453 876	GB_EST14:AA417723 GB_EST11:AA215428 GB_BA1:MTCY77 GB_EST14:AA426336 GB_BA1:STMAACC8	374 303 22255 375 1353	AA417723 AA215428 Z95389 AA426336 M55426	2v01b12.s1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:746207 3' slmilar to contains Alu repetitive element; contains element L1 repetitive element; mRNA sequence. zr95a07.s1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:683412 3' similar to contains Alu repetitive element; mRNA sequence. Mycobacterium tuberculosis H37Rv complete genome; segment 146/162. zv53g02.s1 Soares_testis_NHT Homo sapiens cDNA clone IMAGE:757394 3'; mRNA sequence. S.fradiae aminoglycoside acetyltransferase (aacC8) gene, complete cds.	Homo sapiens Homo sapiens Mycobacterium tuberculosis Homo sapiens Streptomyces fradiae	38,770 39,934 38,889 38,043 37,097	16-OCT-1997 13-Aug-97 18-Jun-98 16-OCT-1997 05-MAY-1993
ка02474 897 ка02480 1779	GB_BA1:AB009078 GB_OM:BTU71200 GB_EST2:F12685 GB_BA1:MTV012	77538 2686 877 287 70287	AC004500 AB009078 U71200 F12685 AL021287	Homo sapiens chromosome 5, P1 clone 1076B9 (LBNL H14), complete sequence. Brevibacterium saccharolyticum gene for L-2.3-butanediol dehydrogenase, complete cds. Bos taurus acetoin reductase mRNA, complete cds. HSC3DA031 normalized infant brain cDNA Homo sapiens cDNA clone c-3da03, mRNA sequence Mycobacterium tuberculosis H37Rv complete genome; segment 132/162.	Homo sapiens Brevibacterium saccharolyticum Bos taurus Homo sapiens Mycobacterium tuberculosis	33,256 96,990 51,659 41,509 36,737	30-MAR-1998 13-Feb-99 8-Oct-97 14-Mar-95 23-Jun-99

WO 01/00844

	GB_BA1:SC6G10	36734	AL049497	Table 4 (continued) Streptomyces coelicolor cosmid 6G10.	Streptomyces coelicolor	35,511	24-MAR-1999	
rxa02485	GB_BA1:AP000060	347800	AP000060	347800 AP000060 Aeropyrum pernix genomic DNA, section 3/7.	Aeropyrum pernix	48,014	22-Jun-99	
				ورد د مند				
rxa02492 840	GB_BA1:STMPGM	921	M83661	Streptomyces coelicolor phosphoglycerate mutase (PGM) gene, complete cds.	Streptomyces coelicolor	65,672	26-Apr-93	
	GB_BA1:MTCY20G9	37218	Z77162	Mycobacterium tuberculosis H37Rv complete genome; segment 25/162.	Mycobacterium tuberculosis	61,436	17-Jun-98	
	GB_BA1:U00018	42991	U00018	Mycobacterium leprae cosmid B2168.	Mycobacterium leprae	37,893	01-MAR-1994	
rxa02528 1098	GB_PR2:HS161N10	56075	AL008707	Human DNA sequence from PAC 161N10 on chromosome Xq25. Contains EST.	Homo sapiens	37,051	23-Nov-99	
	GB_HTG2:AC008235	136017	136017 AC008235		Drosophila melanogaster	36,822	2-Aug-99	
	GB_HTG2:AC008235		136017 AC008235		Drosophila melanogaster	36,822	2-Aug-99	
rxa02539 1641	GB_BA2:RSU17129	17425	U17129	Rhodococcus erythropolis ThcA (thcA) gene, complete cds; and unknown genes.	Rhodococcus erythropolis	66,117	16-Jul-99	•
	GB_BA1:MTV038	16094	AL021933	Mycobacterium tuberculosis H37Rv complete genome; segment 24/162.	Mycobacterium tuberculosis	65,174	17-Jun-98	
	GB_BA2:AF068264	3152	AF068264	Pseudomonas aeruginosa quinoprotein ethanol dehydrogenase (exaA)gene, partial cds; cytochrome c550 precursor (exaB), NAD+ dependent acetaldehyde dehydrogenase (exaC), and pyrroloquinoline quinone synthesis A (pqqA) genes, complete cds; and pyrroloquinoline quinone synthesis B (pqqB) gene, partial cds.		65,448	18-MAR-1999	
rxa02551 483	GB_BA1:BACHYPTP	17057	D29985	Bacillus subtilis wapA and orf genes for wall-associated protein and hypothetical proteins.	Bacillus subtilis	53,602	7-Feb-99	
	GB_BA1:BACHUTWAPÆ8954 GB_BA1:BSGBGLUC 4290	7/28954 4290	D31856 Z34526	Bacillus subtilis genome containing the hut and wapA loci. B.subtilis (Marburg 168) genes for beta-glucoside permease and beta-glucosidase	Bacillus subtilis Bacillus subtilis	53,602 53,602	7-Feb-99 3-Jul-95	_
xa02556 1281	GB_HTG3:AC008128 GB_HTG3:AC008128 GB_PL2:AC005292	335761 335761 99053	AC008128 AC008128 AC005292	groomers appears, *** SEQUENCING IN PROGRESS ***, 106 unordered pieces. Homo sapiens, *** SEQUENCING IN PROGRESS ***, 106 unordered pieces. Genomic sequence for Arabidopsis thaliana BAC F26F24, complete sequence.	Homo sapiens Homo sapiens Arabidopsis thaliana	34,022 34,022 33,858	22-Aug-99 22-Aug-99 16-Apr-99	
rxa02560 990	GB_EST32:AI731605	35692 566	Z66511 Al731605	Caenorhabditis elegans cosmid F07A11, complete sequence. BNLGHi10201 Six-day Cotton fiber Gossypium hirsutum cDNA 5' similar to (AC004684) hypothetical protein [Arabidopsis thaliana], mRNA sequence.	Caenorhabditis elegans Gossypium hirsutum	36,420	2-Sep-99 11-Jun-99	
	GB_INT:CEFU/A11	35692	266511	Caenorhabditis elegans cosmid F07A11, complete sequence.	Caenorhabditis elegans	33,707	2-Sep-99	

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WO 01/00844

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	17-Jun-98	17-Jun-98	29-Sep-99	24-Jun-99	30-OCT-1998	25-MAR-1998	17-Jun-98	10-DEC-1996	29-Sep-94	17-Jun-98	10-DEC-1996	25-MAY-1995	12-Aug-99		86-Jac-2	24-Jun-99	18-MAY-1997	28-Aug-99 5-Jan-98	5-Aug-99	13-Feb-97	01-MAY-1998
	61,677	37,170	19,820	36,957	67,627	70,417	38,532	60,575	57,486	38,018	58,510	57,193	36,858		50,667 50,667	39,187	59,273	58,339 39,637	33,735	35,431	38,851
	Mycobacterium	Mycobacterium tuberculosis	Homo sapiens	Mycobacterium fuberculosis	Mycobacterium	Mycobacterium	tubercutosis Mycobacterium tubercutosis	Mycobacterium tuberculosis	Mycobacterium leprae	Mycobacterium tuberculosis	Mycobacterium tuberculosis		aureofaciens Corynebacterium	glutamicum	Caeronnabolis eregans - Rattus norvegicus	Mycobacterium tuberculosis	Pseudomonas aeruginosa	Pseudomonas tolaasii Arabidopsis thaliana	Arabidopsis thaliana	Mus musculus	Homo sapiens
Table 4 (continued)	Mycobacterium tuberculosis H37Rv complete genome; segment 16/162.	Mycobacterium tuberculosis H37Rv complete genome; segment 16/162.	Homo sapiens chromosome 21 clone LLNLc116H0124 map 21q21, *** SEQUENCING IN PROGRESS ***, in unordered pieces.	Mycobacterium tuberculosis H37Rv complete genome; segment 157/162.	Mycobacterium tuberculosis UDP-galactopyranose mutase (glf) gene, complete Mycobacterium	Mycobacterium tuberculosis UDP-galactopyranose mutase (glf) gene, complete	cus. Mycobacterium tuberculosis H37Rv complete genome; segment 59/162.	Mycobacterium tuberculosis sequence from clone y151.	Mycobacterium leprae cosmid B1549.	Mycobacterium tuberculosis H3/Rv complete genome; segment 59/162.	Mycobacterium tuberculosis sequence from clone y151.	Streptomyces aureofaciens glycogen branching enzyme (glgB) gene, complete	cds. Corynebacterium glutamicum yjcc gene, amtR gene and citE gene, partial.	constitute delument 201M Limons annuals sililadadas de constituto de con	Odenioniabunis dregaris Cosmu in rod, Comprete sequence. UI-R-C3-sz-h-03-0-UI-s1 UI-R-C3 Rattus norvegicus cDNA clone UI-R-C3-sz-h- Rattus norvegicus 03-0-UI-31 mRNA sequence.	Mycobacterium tuberculosis H37Rv complete genome; segment 155/162.	Pseudomonas aeruginosa (orfX), glycerol dfiffusion facilitator (glpF), glycerol kinase (glpK), and Glp repressor (glpR) genes, complete cds, and (orfK) gene, partial cds.		-	sequence. Was musculus Btk locus, alpha-D-galactosidase A (Ags), ribosomal protein	(L44L), and bruton's tyrosme kinase (bit) genes, complete cds. Homo sapiens chromosome 16, cosmid clone 363E3 (LANL), complete sequence.
	Z96800	Z96800	AL121632	AL022076	AF026540	U96128	Z73902	AD000018	U00014	Z73902	AD000018	L11647	AJ133719	746036	A1547662	AL022121	U49666	AB015974 N65787	AC005916	U58105	AC004643
	38900	38900	46989	23740	1778	1200	32514	37036	36470	32514	37036	2557	1839	30073	377	121125	4495	1641 512	65839	88871	43411
	GB_BA1:MTCY63	GB_BA1:MTCY63	GB_HTG1:HS24H01	GB_BA1:MTV026	GB_BA2:AF026540	GB_BA2:MTU96128	GB_BA1:MTCY130	GB_BA1:MSGY151	GB_BA1:U00014	GB_BA1:MTCY130	GB_BA1:MSGY151	GB_BA1:STMGLGEN	GB_BA1:CGL133719	SOLVED AND ASSESSMENT	GB_EST29:AI547662	GB_BA1:MTV025	GB_BA1:PAU49666	GB_BA1:AB015974 GB_EST6:N65787	GB_PL2:T17H3	GB_RO:MMU58105	GB_PR3:AC004643
	rxa02572 668			rxa02596 1326			rxa02611 1775			rxa02612 2316			rxa02621 942			rxa02640 1650		rxa02654 1008			rxa02666 891

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01-MAY-1998	1-Jul-98	20-MAY-1993	17-Jun-98	27-Jan-94 26-Apr-93 26-Apr-93	29-Jul-93	27-Jun-97	27-Jun-97	29-DEC-1998	27-Jun-97	15-Nov-99 24-Jun-98	19-Jun-98	27-Jul-98 24-Jun-98	12-Jul-99 27-Jul-98 20-Feb-99
41,599	40,413	40,735	36,471	38,477 57,371 57,277	57,277	50,746	36,364	37,059	42,149	37,655 99,580	38,363	39,444 98,226	60,399 36,426 99,640
Homo sapiens	Corynebacterium glutamicum	Paracoccus denitrificans	Mycobacterium	Myxococcus xanthus Bacillus caldolyticus Bacillus	stearothermophilus Bacillus	stearothermophilus Danio rerio	Danio rerio	A Homo sapiens	Danio rerio	Homo sapiens Corynebacterium	glutamicum Mycobacterium	tuberculosis Streptomyces coelicolor Corynebacterium	glutamicum Streptomyces coelicolor Streptomyces coelicolor Corynebacterium glutamicum
Table 4 (continued) Homo sapiens chromosome 16, cosmid clone 363E3 (LANL), complete	sequence. Corynebacterium glutamicum N-acetylglutamylphosphate reductase (argC), ornithine acetyltransferase (argJ), N-acetylglutamate kinase (argB), acetylornithine transaminase (argD), ornithine carbamoytransferase (argF)	arginine repressor (argR), argininosuccinate synthase (argG), and argininosuccinate lyase (argH) genes, complete cds. Paracoccus denitrificans NADH dehydrogenase (URF4), (NQO8), (NQO9), (URF5), (URF6), (NQO10), (NQO11), (NQO12), (NQO13), and (NQO14) genes, complete cds's; biotin [acetyl-CoA carboxyl] ligase (birA) gene, complete	cos. Mycobacterium tuberculosis H37Rv complete genome; segment 101/162.	Myxococcus xanthus devR and devS genes, complete cds's. B.caldolyticus lactate dehydrogenase (LDH) gene, complete cds. B.stearothermophilus lct gene encoding L-lactate dehydrogenase, complete	cds. B.stearothermophilus lct gene.	fa09d04.r1 Zebrafish ICRFzfls Danio rerio cDNA clone 11A22 5' similar to TR:G1171163 G1171163 G7-MISMATCH BINDING PROTEIN :; mRNA	sequence. fa09d04.r1 Zebrafish ICRFzfls Danio rerio cDNA clone 11A22 5' similar to TR:G1171163 G1171163 G/T-MISMATCH BINDING PROTEIN.; mRNA sequence.	ah67d06.s1 Soares_testis_NHT Homo sapiens cDNA clone 1320683 3', mRNA Homo sapiens	sequence. fa09d04.r1 Zebrafish ICRFzfls Danio rerio cDNA clone 11A22 5' similar to TR:G1171163 G1171163 G/T-MISMATCH BINDING PROTEIN :; mRNA	sequence. Homo sapiens, complete sequence. gDNA encoding glucose-6-phosphate dehydrogenase.	Mycobacterium tuberculosis H37Rv complete genome; segment 63/162.	Streptomyces coelicolor cosmid 5A7. gDNA encoding glucose-6-phosphate dehydrogenase.	Streptomyces coelicolor cosmid C22. Streptomyces coelicolor cosmid 5A7. Corynebacterium glutamicum tkt gene for transketolase, complete cds.
AC004643	AF049897	L02354	Z 77163	L19029 M19394 M14788	A06664	AA494626	AA494626	AA758660	AA494626	150172 AC006285 2260 E13655	Z 95844	AL031107 E13655	AL096839 AL031107 AB023377
43411	9196	10425	42861	2452 1147 1361	1350	121	121	233	121	150172 2260	40790	40337 2260	22115 40337 2572
GB_PR3:AC004643	GB_BA2:AF049897	GB_BA1:PDENQOURF 10425	GB_BA1:MTCY339	GB_BA1:MXADEVRS GB_BA1:BACLDH GB_BA1:BACLDHL	GB_PAT:A06664	GB_EST15:AA494626	GB_EST15:AA494626	GB_EST19:AA758660	GB_EST15:AA494626	GB_PR4:AC006285 GB_PAT:E13655	GB_BA1:MTCY493	GB_BA1:SC5A7 GB_PAT:E13655	GB_BA1:SCC22 GB_BA1:SC5A7 GB_BA1:AB023377
		אמס2675 1980		ка02694 1065		rxa02729 844		rxa02730 1161		rxa02737 1665		rxa02738 1203	rxa02739 2223

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04-DEC-1998 01-MAR-1994	2-Aug-99	2-Aug-99	20-Sep-99	12-Jun-98	12-Jun-98	23-Jun-99	04-DEC-1998 01-MAR-1994		26-Jun-99	8-Sep-97	08-OCT-1998	1-Feb-97	01-MAR-1994	9-Apr-97	20-Aug-98	66-IDC-27	22-Jul-99	30-Jun-99
61,573 61,573	37,105	37,105	38,728	33,116	33,116	36,379	48,401 48,401		37,128	38,889	34,321	38,072	34,462	. 50,445	59,314	100'10	37,607	40,157
Mycobacterium leprae Mycobacterium leprae	Drosophila melanogaster	Drosophila melanogaster	Orosophila melanogaster	Homo sapiens	Homo sapiens	Ephydatia fluviatilis	Mycobacterium leprae Mycobacterium leprae		Homo sapiens	Corynebacterium	giutamicum Homo sapiens	Bacillus firmus	Mycobacterium leprae	Pseudomonas syringae pv. 50,445 syringae	Streptomyces coelicolor	nomo sapiens	Homo sapiens	e Mus musculus
Table 4 (continued) Mycobacterium leprae cosmid L536. Mycobacterium leprae cosmid B1496.	Drosophila melanogaster chromosome 2 clone BACR48110 (D505) RPCI-98 48.I.10 map 49E6-49F8 strain y; cn bw sp, *** SEQUENCING IN PROGRESS *** 17 unordered pieces.	Drosophila melangaster chromosome 2 clone BACR48I10 (D505) RPCI-98 48.I.10 may 49E6-49F8 strain y; cn bw sp, *** SEQUENCING IN PROGRESS *** 17 unordered pieces	Drosophila melanogaster chromosome 2 clone BACR16P13 (D597) RPCI-98 16.P.13 map 49E-49F strain y; cn bw sp, *** SEQUENCING IN PROGRESS*** 87 unordered pieces.	Homo sapiens clone DJ1022114, *** SEQUENCING IN PROGRESS ***, 14	Homo sapiens clone DJ1022I14, *** SEQUENCING IN PROGRESS ***, 14 unordered pieces.	Ephydatia fluviatilis mRNA for G protein a subunit 4, partial cds.	Mycobacterium leprae cosmid L536. Mycobacterium leprae cosmid B1496.		Homo sapiens clone NH0501007, *** SEQUENCING IN PROGRESS ***, 3 unordered pieces.	C.glutamicum betP gene.	glutamicum HS_3136_A1_A03_MR CIT Approved Human Genomic Sperm Library D Homo Homo sapiens sapiens genomic clone Plate=3136 Col=5 Row=A, genomic survey sequence.	Bacillus firmus dppABC operon, dipeptide transporter protein dppA gene, partial cds, and dipeptide transporter proteins dppB and dppC genes, complete cds.	Mycobacterium leprae cosmid B229.	Pseudomonas syringae pv. syringae putative dihydropteroate synthase gene, partial cds, regulatory protein MrsA (mrsA), triose phosphate isomerase (tpiA), transport protein SecG (secG), tRNA-Leu, tRNA-Met, and 15 kDa protein genes, complete cds.		Normo sapiens chiginosome 17 clone 2020_N_17 map 17, SEQUENCING IN PROGRESS ***, 12 unordered pieces.		IN FROOKESSS 1, 12 unoutered pieces. AV117143 Mus musculus C57BL/6J 10-day embryo Mus musculus cDNA clone Mus musculus 2610200J17, mRNA sequence.
Z99125 U00013	174368 AC006247	174368 AC006247	121474 AC007150	129429 AC004951	129429 AC004951	AB006546	Z99125 U00013		AC007401	X93514	AQ148714	U64514	020000	U85643	AL031317	AC008103	AC008105	AV117143
36224 35881	174368	174368	121474	129429	129429	931	36224 35881		83657	2339	405	3837	36947	4032	41055	91421	91421	222
GB_BA1:MLCL536 GB_BA1:U00013	GB_HTG2:AC006247	GB_HTG2:AC006247	GB_HTG3:AC007150	GB_HTG2:AC004951	GB_HTG2:AC004951	GB_IN1:AB006546	GB_BA1:MLCL536 GB_BA1:U00013		GB_HTG2:AC007401	GB_BA1:CGBETPGEN 2339	GB_GSS9:AQ148714	GB_BA1:BFU64514	GB_BA1:U00020	GB_BA2:PSU85643	GB_BA1:SC6G4	cb_di_62:Acousting	GB_HTG2:AC008105	GB_EST33:AV117143
	א 1053 אמי			rxa02741 1089			rxa02743 1161			rxa02797 1026			xa02803 680			rxaU2621 363		

WO 01/00844

				[able 4 (continued)			
rxa02829 373	GB_HTG1:HSU9G8	48735	AL008714	48735 AL008714 Homo sapiens chromosome X clone LL0XNC01-9G8, *** SEQUENCING IN	Homo sapiens	41,595	23-Nov-99
	GB_HTG1:HSU9G8	48735	48735 AL008714	PROGRESS ***, in unordered pieces. Home sabiens chromosome X clone 11.0XNC01-9G8 *** SFOLIENCING IN	Homo e o cione	11 505	
	ı			PROGRESS ***, in unordered pieces.		660'- +	66-40N-67
	GB_PR3:HSU85B5	39550	Z69724	Human DNA sequence from cosmid U85B5, between markers DXS366 and	Homo sapiens	41.595	23-Nov-99
				DXS87 on chromosome X.	•		
xc03216 1141		151720	AC008184	GB_HTG3:AC008184 151720 AC008184 Drosophila melanogaster chromosome 2 clone BACR04D05 (D540) RPCI-98	Drosophila melanogaster	39.600	2-Aug-99
				04.D.5 map 36E5-36F2 strain y; cn bw sp, *** SEQUENCING IN PROGRESS			n 1
				***, 27 unordered pieces.			
	GB_EST15:AA477537 411	411	AA477537	zu36g12.r1 Soares ovary tumor NbHOT Homo sapiens cDNA clone	Homo sapiens	37.260	9-Nov-97
				IMAGE:740134 5' similar to contains Alu repetitive element contains element			
				HGR repetitive element ;, mRNA sequence.			
	GB_EST26:AI330662	412	AI330662	fa91d08.y1 zebrafish fin day1 regeneration Danio rerio cDNA 5', mRNA	Danio rerio	37,805	28-DFC-1998
				sequence.			
rxs03215 1038	GB_BA1:SC3F9	19830	19830 AL023862	Streptomyces coelicolor cosmid 3F9.	Streptomyces coelicolor	48,657	10-Feb-99
					A3(2)		
	GB_BA1:SLLINC	36270	X79146	S.lincolnensis (78-11) Lincomycin production genes.	Streptomyces lincolnensis	39,430	15-MAY-1996
	GB HTG5:AC009660	204320	204320 AC009660 Homo				
				IN PROGRESS ***, 41 unordered pieces.	romo sapiens	35,151	04-DEC-1999
rxs03224 1288	GB_PR3:AC004076	41322	41322 AC004076 Homo	Homo sapiens chromosome 19, cosmid R30217, complete sequence.	Homo sapiens	37.788	29-Jan-98
•	GB_PL2:SPAC926	23193	AL110469	S.pombe chromosome I cosmid c926.	Schizosaccharomyces	38,474	2-Sep-99
	GB_BA2:AE001081	11473	AE001081	AE001081 Archaeoglobus fulgidus section 26 of 172 of the complete genome.	pombe Archaeoglobus fulgidus	35.871	15-DEC-1997

Exemplification

Example 1: Preparation of total genomic DNA of Corynebacterium glutamicum ATCC 13032

A culture of Corynebacterium glutamicum (ATCC 13032) was grown overnight 5 at 30°C with vigorous shaking in BHI medium (Difco). The cells were harvested by centrifugation, the supernatant was discarded and the cells were resuspended in 5 ml buffer-I (5% of the original volume of the culture — all indicated volumes have been calculated for 100 ml of culture volume). Composition of buffer-I: 140.34 g/l sucrose, 2.46 g/l MgSO₄ x 7H₂O₅ 10 ml/l KH₂PO₄ solution (100 g/l, adjusted to pH 6.7 with KOH), 50 ml/l M12 concentrate (10 g/l (NH₄)₂SO₄, 1 g/l NaCl, 2 g/l MgSO₄ x 7H₂O, 0.2 g/l CaCl₂, 0.5 g/l yeast extract (Difco), 10 ml/l trace-elements-mix (200 mg/l FeSO₄ x H₂O, 10 mg/l ZnSO₄ x 7 H₂O, 3 mg/l MnCl₂ x 4 H₂O, 30 mg/l H₃BO₃ 20 mg/l CoCl₂ x 6 H₂O, 1 mg/l NiCl₂ x 6 H₂O, 3 mg/l Na₂MoO₄ x 2 H₂O, 500 mg/l complexing agent 15 (EDTA or critic acid), 100 ml/l vitamins-mix (0.2 mg/l biotin, 0.2 mg/l folic acid, 20 mg/l p-amino benzoic acid, 20 mg/l riboflavin, 40 mg/l ca-panthothenate, 140 mg/l nicotinic acid, 40 mg/l pyridoxole hydrochloride, 200 mg/l myo-inositol). Lysozyme was added to the suspension to a final concentration of 2.5 mg/ml. After an approximately 4 h incubation at 37°C, the cell wall was degraded and the resulting protoplasts are harvested by centrifugation. The pellet was washed once with 5 ml 20 buffer-I and once with 5 ml TE-buffer (10 mM Tris-HCl, I mM EDTA, pH 8). The pellet was resuspended in 4 ml TE-buffer and 0.5 ml SDS solution (10%) and 0.5 ml NaCl solution (5 M) are added. After adding of proteinase K to a final concentration of 200 µg/ml, the suspension is incubated for ca.18 h at 37°C. The DNA was purified by extraction with phenol, phenol-chloroform-isoamylalcohol and chloroform-25 isoamylalcohol using standard procedures. Then, the DNA was precipitated by adding 1/50 volume of 3 M sodium acetate and 2 volumes of ethanol, followed by a 30 min incubation at -20°C and a 30 min centrifugation at 12,000 rpm in a high speed centrifuge using a SS34 rotor (Sorvall). The DNA was dissolved in 1 ml TE-buffer containing 20 μg/ml RNaseA and dialysed at 4°C against 1000 ml TE-buffer for at least 3 hours. During this time, the buffer was exchanged 3 times. To aliquots of 0.4 ml of the dialysed DNA solution, 0.4 ml of 2 M LiCl and 0.8 ml of ethanol are added. After a 30

- 119 -

min incubation at -20°C, the DNA was collected by centrifugation (13,000 rpm, Biofuge Fresco, Heraeus, Hanau, Germany). The DNA pellet was dissolved in TE-buffer. DNA prepared by this procedure could be used for all purposes, including southern blotting or construction of genomic libraries.

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Example 2: Construction of genomic libraries in *Escherichia coli* of *Corynebacterium glutamicum* ATCC13032.

Using DNA prepared as described in Example 1, cosmid and plasmid libraries were constructed according to known and well established methods (see e.g., Sambrook, J. et al. (1989) "Molecular Cloning: A Laboratory Manual", Cold Spring Harbor Laboratory Press, or Ausubel, F.M. et al. (1994) "Current Protocols in Molecular Biology", John Wiley & Sons.)

Any plasmid or cosmid could be used. Of particular use were the plasmids pBR322 (Sutcliffe, J.G. (1979) *Proc. Natl. Acad. Sci. USA*, 75:3737-3741); pACYC177 (Change & Cohen (1978) *J. Bacteriol* 134:1141-1156), plasmids of the pBS series (pBSSK+, pBSSK- and others; Stratagene, LaJolla, USA), or cosmids as SuperCos1 (Stratagene, LaJolla, USA) or Lorist6 (Gibson, T.J., Rosenthal A. and Waterson, R.H. (1987) *Gene* 53:283-286. Gene libraries specifically for use in *C. glutamicum* may be constructed using plasmid pSL109 (Lee, H.-S. and A. J. Sinskey (1994) *J. Microbiol. Biotechnol.* 4: 256-263).

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Example 3: DNA Sequencing and Computational Functional Analysis

Genomic libraries as described in Example 2 were used for DNA sequencing according to standard methods, in particular by the chain termination method using ABI377 sequencing machines (see *e.g.*, Fleischman, R.D. *et al.* (1995) "Whole-genome Random Sequencing and Assembly of Haemophilus Influenzae Rd., *Science*, 269:496-512). Sequencing primers with the following nucleotide sequences were used: 5'-GGAAACAGTATGACCATG-3' or 5'-GTAAAACGACGCCCAGT-3'.

Example 4: In vivo Mutagenesis

30 In vivo mutagenesis of Corynebacterium glutamicum can be performed by passage of plasmid (or other vector) DNA through E. coli or other microorganisms (e.g. Bacillus spp. or yeasts such as Saccharomyces cerevisiae) which are impaired in their capabilities to maintain

the integrity of their genetic information. Typical mutator strains have mutations in the genes for the DNA repair system (e.g., mutHLS, mutD, mutT, etc.; for reference, see Rupp, W.D. (1996) DNA repair mechanisms, in: *Escherichia coli* and *Salmonella*, p. 2277-2294, ASM: Washington.) Such strains are well known to those of ordinary skill in the art. The use of such strains is illustrated, for example, in Greener, A. and Callahan, M. (1994) *Strategies* 7: 32-34.

Example 5: DNA Transfer Between Escherichia coli and Corynebacterium glutamicum

Several Corynebacterium and Brevibacterium species contain endogenous plasmids (as e.g., pHM1519 or pBL1) which replicate autonomously (for review see, e.g., Martin, J.F. et al. (1987) Biotechnology, 5:137-146). Shuttle vectors for Escherichia coli and Corynebacterium glutamicum can be readily constructed by using standard vectors for E. coli (Sambrook, J. et al. (1989), "Molecular Cloning: A Laboratory Manual", Cold Spring Harbor Laboratory Press or Ausubel, F.M. et al. (1994) "Current Protocols in 15 Molecular Biology", John Wiley & Sons) to which a origin or replication for and a suitable marker from Corynebacterium glutamicum is added. Such origins of replication are preferably taken from endogenous plasmids isolated from Corynebacterium and Brevibacterium species. Of particular use as transformation markers for these species are genes for kanamycin resistance (such as those derived from the Tn5 or Tn903 transposons) or chloramphenicol (Winnacker, E.L. (1987) "From Genes to Clones — 20 Introduction to Gene Technology, VCH, Weinheim). There are numerous examples in the literature of the construction of a wide variety of shuttle vectors which replicate in both E. coli and C. glutamicum, and which can be used for several purposes, including gene overexpression (for reference, see e.g., Yoshihama, M. et al. (1985) J. Bacteriol. 162:591-597, 25 Martin J.F. et al. (1987) Biotechnology, 5:137-146 and Eikmanns, B.J. et al. (1991) Gene, 102:93-98).

Using standard methods, it is possible to clone a gene of interest into one of the shuffle vectors described above and to introduce such a hybrid vectors into strains of Corynebacterium glutamicum. Transformation of C. glutamicum can be achieved by protoplast transformation (Kastsumata, R. et al. (1984) J. Bacteriol. 159306-311), electroporation (Liebl, E. et al. (1989) FEMS Microbiol. Letters, 53:399-303) and in cases where special vectors are used, also by conjugation (as described e.g. in Schäfer, A et al.

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(1990) J. Bacteriol. 172:1663-1666). It is also possible to transfer the shuttle vectors for C. glutamicum to E. coli by preparing plasmid DNA from C. glutamicum (using standard methods well-known in the art) and transforming it into E. coli. This transformation step can be performed using standard methods, but it is advantageous to use an Mcr-deficient E. coli strain, such as NM522 (Gough & Murray (1983) J. Mol. Biol. 166:1-19).

Genes may be overexpressed in *C. glutamicum* strains using plasmids which comprise pCG1 (U.S. Patent No. 4,617,267) or fragments thereof, and optionally the gene for kanamycin resistance from TN903 (Grindley, N.D. and Joyce, C.M. (1980) *Proc. Natl. Acad. Sci. USA* 77(12): 7176-7180). In addition, genes may be overexpressed in *C. glutamicum* strains using plasmid pSL109 (Lee, H.-S. and A. J. Sinskey (1994) *J. Microbiol. Biotechnol.* 4: 256-263).

Aside from the use of replicative plasmids, gene overexpression can also be achieved by integration into the genome. Genomic integration in *C. glutamicum* or other Corynebacterium or Brevibacterium species may be accomplished by well-known methods, such as homologous recombination with genomic region(s), restriction endonuclease mediated integration (REMI) (see, *e.g.*, DE Patent 19823834), or through the use of transposons. It is also possible to modulate the activity of a gene of interest by modifying the regulatory regions (*e.g.*, a promoter, a repressor, and/or an enhancer) by sequence modification, insertion, or deletion using site-directed methods (such as homologous recombination) or methods based on random events (such as transposon mutagenesis or REMI). Nucleic acid sequences which function as transcriptional terminators may also be inserted 3' to the coding region of one or more genes of the invention; such terminators are well-known in the art and are described, for example, in Winnacker, E.L. (1987) From Genes to Clones – Introduction to Gene Technology. VCH: Weinheim.

Example 6: Assessment of the Expression of the Mutant Protein

Observations of the activity of a mutated protein in a transformed host cell rely on the fact that the mutant protein is expressed in a similar fashion and in a similar quantity to that of the wild-type protein. A useful method to ascertain the level of transcription of the mutant gene (an indicator of the amount of mRNA available for translation to the gene product) is to perform a Northern blot (for reference see, for example, Ausubel *et al.*

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(1988) Current Protocols in Molecular Biology, Wiley: New York), in which a primer designed to bind to the gene of interest is labeled with a detectable tag (usually radioactive or chemiluminescent), such that when the total RNA of a culture of the organism is extracted, run on gel, transferred to a stable matrix and incubated with this probe, the binding and quantity of binding of the probe indicates the presence and also the quantity of mRNA for this gene. This information is evidence of the degree of transcription of the mutant gene. Total cellular RNA can be prepared from *Corynebacterium glutamicum* by several methods, all well-known in the art, such as that described in Bormann, E.R. et al. (1992) *Mol. Microbiol.* 6: 317-326.

To assess the presence or relative quantity of protein translated from this mRNA, standard techniques, such as a Western blot, may be employed (see, for example, Ausubel et al. (1988) Current Protocols in Molecular Biology, Wiley: New York). In this process, total cellular proteins are extracted, separated by gel electrophoresis, transferred to a matrix such as nitrocellulose, and incubated with a probe, such as an antibody, which specifically binds to the desired protein. This probe is generally tagged with a chemiluminescent or colorimetric label which may be readily detected. The presence and quantity of label observed indicates the presence and quantity of the desired mutant protein present in the cell.

20 Example 7: Growth of Genetically Modified *Corynebacterium glutamicum* — Media and Culture Conditions

Genetically modified *Corynebacteria* are cultured in synthetic or natural growth media. A number of different growth media for Corynebacteria are both well-known and readily available (Lieb *et al.* (1989) *Appl. Microbiol. Biotechnol.*, 32:205-210; von der Osten *et al.* (1998) *Biotechnology Letters*, 11:11-16; Patent DE 4,120,867; Liebl (1992) "The Genus *Corynebacterium*, in: The Procaryotes, Volume II, Balows, A. *et al.*, eds. Springer-Verlag). These media consist of one or more carbon sources, nitrogen sources, inorganic salts, vitamins and trace elements. Preferred carbon sources are sugars, such as mono-, di-, or polysaccharides. For example, glucose, fructose, mannose, galactose, ribose, sorbose, ribulose, lactose, maltose, sucrose, raffinose, starch or cellulose serve as very good carbon sources. It is also possible to supply sugar to the media via complex compounds such as molasses or other by-products from sugar refinement. It can also be

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advantageous to supply mixtures of different carbon sources. Other possible carbon sources are alcohols and organic acids, such as methanol, ethanol, acetic acid or lactic acid. Nitrogen sources are usually organic or inorganic nitrogen compounds, or materials which contain these compounds. Exemplary nitrogen sources include ammonia gas or ammonia salts, such as NH₄Cl or (NH₄)₂SO₄, NH₄OH, nitrates, urea, amino acids or complex nitrogen sources like corn steep liquor, soy bean flour, soy bean protein, yeast extract, meat extract and others.

Inorganic salt compounds which may be included in the media include the chloride-, phosphorous- or sulfate- salts of calcium, magnesium, sodium, cobalt, molybdenum, potassium, manganese, zinc, copper and iron. Chelating compounds can be added to the medium to keep the metal ions in solution. Particularly useful chelating compounds include dihydroxyphenols, like catechol or protocatechuate, or organic acids, such as citric acid. It is typical for the media to also contain other growth factors, such as vitamins or growth promoters, examples of which include biotin, riboflavin, thiamin, folic acid, nicotinic acid, pantothenate and pyridoxin. Growth factors and salts frequently originate from complex media components such as yeast extract, molasses, corn steep liquor and others. The exact composition of the media compounds depends strongly on the immediate experiment and is individually decided for each specific case. Information about media optimization is available in the textbook "Applied Microbiol. Physiology, A Practical Approach (eds. P.M. Rhodes, P.F. Stanbury, IRL Press (1997) pp. 53-73, ISBN 0 19 963577 3). It is also possible to select growth media from commercial suppliers, like standard 1 (Merck) or BHI (grain heart infusion, DIFCO) or others.

All medium components are sterilized, either by heat (20 minutes at 1.5 bar and 121°C) or by sterile filtration. The components can either be sterilized together or, if necessary, separately. All media components can be present at the beginning of growth, or they can optionally be added continuously or batchwise.

Culture conditions are defined separately for each experiment. The temperature should be in a range between 15°C and 45°C. The temperature can be kept constant or can be altered during the experiment. The pH of the medium should be in the range of 5 to 8.5, preferably around 7.0, and can be maintained by the addition of buffers to the media. An exemplary buffer for this purpose is a potassium phosphate buffer. Synthetic buffers such as MOPS, HEPES, ACES and others can alternatively or simultaneously be used. It

is also possible to maintain a constant culture pH through the addition of NaOH or NH₄OH during growth. If complex medium components such as yeast extract are utilized, the necessity for additional buffers may be reduced, due to the fact that many complex compounds have high buffer capacities. If a fermentor is utilized for culturing the microorganisms, the pH can also be controlled using gaseous ammonia.

The incubation time is usually in a range from several hours to several days. This time is selected in order to permit the maximal amount of product to accumulate in the broth. The disclosed growth experiments can be carried out in a variety of vessels, such as microtiter plates, glass tubes, glass flasks or glass or metal fermentors of different sizes. For screening a large number of clones, the microorganisms should be cultured in microtiter plates, glass tubes or shake flasks, either with or without baffles. Preferably 100 ml shake flasks are used, filled with 10% (by volume) of the required growth medium. The flasks should be shaken on a rotary shaker (amplitude 25 mm) using a speed-range of 100 - 300 rpm. Evaporation losses can be diminished by the maintenance of a humid atmosphere; alternatively, a mathematical correction for evaporation losses should be performed.

If genetically modified clones are tested, an unmodified control clone or a control clone containing the basic plasmid without any insert should also be tested. The medium is inoculated to an OD₆₀₀ of O.5 – 1.5 using cells grown on agar plates, such as CM plates (10 g/l glucose, 2,5 g/l NaCl, 2 g/l urea, 10 g/l polypeptone, 5 g/l yeast extract, 5 g/l meat extract, 22 g/l NaCl, 2 g/l urea, 10 g/l polypeptone, 5 g/l yeast extract, 5 g/l meat extract, 22 g/l agar, pH 6.8 with 2M NaOH) that had been incubated at 30°C. Inoculation of the media is accomplished by either introduction of a saline suspension of *C. glutamicum* cells from CM plates or addition of a liquid preculture of this bacterium.

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Example 8 - In vitro Analysis of the Function of Mutant Proteins

The determination of activities and kinetic parameters of enzymes is well established in the art. Experiments to determine the activity of any given altered enzyme must be tailored to the specific activity of the wild-type enzyme, which is well within the ability of one of ordinary skill in the art. Overviews about enzymes in general, as well as specific details concerning structure, kinetics, principles, methods, applications and examples for the determination of many enzyme activities may be

found, for example, in the following references: Dixon, M., and Webb, E.C., (1979)
Enzymes. Longmans: London; Fersht, (1985) Enzyme Structure and Mechanism.
Freeman: New York; Walsh, (1979) Enzymatic Reaction Mechanisms. Freeman: San Francisco; Price, N.C., Stevens, L. (1982) Fundamentals of Enzymology. Oxford Univ.
Press: Oxford; Boyer, P.D., ed. (1983) The Enzymes, 3rd ed. Academic Press: New York; Bisswanger, H., (1994) Enzymkinetik, 2nd ed. VCH: Weinheim (ISBN 3527300325); Bergmeyer, H.U., Bergmeyer, J., Graßl, M., eds. (1983-1986) Methods of Enzymatic Analysis, 3rd ed., vol. I-XII, Verlag Chemie: Weinheim; and Ullmann's Encyclopedia of Industrial Chemistry (1987) vol. A9, "Enzymes". VCH: Weinheim, p. 352-363.

The activity of proteins which bind to DNA can be measured by several well-established methods, such as DNA band-shift assays (also called gel retardation assays). The effect of such proteins on the expression of other molecules can be measured using reporter gene assays (such as that described in Kolmar, H. et al. (1995) EMBO J. 14: 3895-3904 and references cited therein). Reporter gene test systems are well known and established for applications in both pro- and eukaryotic cells, using enzymes such as beta-galactosidase, green fluorescent protein, and several others.

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The determination of activity of membrane-transport proteins can be performed according to techniques such as those described in Gennis, R.B. (1989) "Pores,

Channels and Transporters", in Biomembranes, Molecular Structure and Function,

Springer: Heidelberg, p. 85-137; 199-234; and 270-322.

Example 9: Analysis of Impact of Mutant Protein on the Production of the Desired Product

The effect of the genetic modification in *C. glutamicum* on production of a desired compound (such as an amino acid) can be assessed by growing the modified microorganism under suitable conditions (such as those described above) and analyzing the medium and/or the cellular component for increased production of the desired product (*i.e.*, an amino acid). Such analysis techniques are well known to one of ordinary skill in the art, and include spectroscopy, thin layer chromatography, staining methods of various kinds, enzymatic and microbiological methods, and analytical chromatography such as high performance liquid chromatography (see, for example,

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Ullman, Encyclopedia of Industrial Chemistry, vol. A2, p. 89-90 and p. 443-613, VCH: Weinheim (1985); Fallon, A. et al., (1987) "Applications of HPLC in Biochemistry" in: Laboratory Techniques in Biochemistry and Molecular Biology, vol. 17; Rehm et al. (1993) Biotechnology, vol. 3, Chapter III: "Product recovery and purification", page 469-714, VCH: Weinheim; Belter, P.A. et al. (1988) Bioseparations: downstream processing for biotechnology, John Wiley and Sons; Kennedy, J.F. and Cabral, J.M.S. (1992) Recovery processes for biological materials, John Wiley and Sons; Shaeiwitz, J.A. and Henry, J.D. (1988) Biochemical separations, in: Ulmann's Encyclopedia of Industrial Chemistry, vol. B3, Chapter 11, page 1-27, VCH: Weinheim; and Dechow,
F.J. (1989) Separation and purification techniques in biotechnology, Noyes Publications.)

In addition to the measurement of the final product of fermentation, it is also possible to analyze other components of the metabolic pathways utilized for the production of the desired compound, such as intermediates and side-products, to determine the overall efficiency of production of the compound. Analysis methods include measurements of nutrient levels in the medium (e.g., sugars, hydrocarbons, nitrogen sources, phosphate, and other ions), measurements of biomass composition and growth, analysis of the production of common metabolites of biosynthetic pathways, and measurement of gasses produced during fermentation. Standard methods for these measurements are outlined in Applied Microbial Physiology, A Practical Approach, P.M. Rhodes and P.F. Stanbury, eds., IRL Press, p. 103-129; 131-163; and 165-192 (ISBN: 0199635773) and references cited therein.

Example 10: Purification of the Desired Product from C. glutamicum Culture

Recovery of the desired product from the *C. glutamicum* cells or supernatant of the above-described culture can be performed by various methods well known in the art. If the desired product is not secreted from the cells, the cells can be harvested from the culture by low-speed centrifugation, the cells can be lysed by standard techniques, such as mechanical force or sonication. The cellular debris is removed by centrifugation, and the supernatant fraction containing the soluble proteins is retained for further purification of the desired compound. If the product is secreted from the *C. glutamicum*

cells, then the cells are removed from the culture by low-speed centrifugation, and the supernate fraction is retained for further purification.

The supernatant fraction from either purification method is subjected to chromatography with a suitable resin, in which the desired molecule is either retained on a chromatography resin while many of the impurities in the sample are not, or where the impurities are retained by the resin while the sample is not. Such chromatography steps may be repeated as necessary, using the same or different chromatography resins. One of ordinary skill in the art would be well-versed in the selection of appropriate chromatography resins and in their most efficacious application for a particular molecule to be purified. The purified product may be concentrated by filtration or ultrafiltration, and stored at a temperature at which the stability of the product is maximized.

There are a wide array of purification methods known to the art and the preceding method of purification is not meant to be limiting. Such purification techniques are described, for example, in Bailey, J.E. & Ollis, D.F. Biochemical Engineering Fundamentals, McGraw-Hill: New York (1986).

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The identity and purity of the isolated compounds may be assessed by techniques standard in the art. These include high-performance liquid chromatography (HPLC), spectroscopic methods, staining methods, thin layer chromatography, NIRS, enzymatic assay, or microbiologically. Such analysis methods are reviewed in: Patek et al. (1994)

20 Appl. Environ. Microbiol. 60: 133-140; Malakhova et al. (1996) Biotekhnologiya 11: 27-32; and Schmidt et al. (1998) Bioprocess Engineer. 19: 67-70. Ulmann's Encyclopedia of Industrial Chemistry, (1996) vol. A27, VCH: Weinheim, p. 89-90, p. 521-540, p. 540-547, p. 559-566, 575-581 and p. 581-587; Michal, G. (1999) Biochemical Pathways: An Atlas of Biochemistry and Molecular Biology, John Wiley and Sons; Fallon, A. et al. (1987) Applications of HPLC in Biochemistry in: Laboratory Techniques in Biochemistry and Molecular Biology, vol. 17.

Example 11: Analysis of the Gene Sequences of the Invention

The comparison of sequences and determination of percent homology between two sequences are art-known techniques, and can be accomplished using a mathematical algorithm, such as the algorithm of Karlin and Altschul (1990) *Proc. Natl. Acad. Sci.* USA 87:2264-68, modified as in Karlin and Altschul (1993) *Proc. Natl. Acad. Sci.* USA

- 128 -

90:5873-77. Such an algorithm is incorporated into the NBLAST and XBLAST programs (version 2.0) of Altschul, *et al.* (1990) *J. Mol. Biol.* 215:403-10. BLAST nucleotide searches can be performed with the NBLAST program, score = 100, wordlength = 12 to obtain nucleotide sequences homologous to SMP nucleic acid molecules of the invention. BLAST protein searches can be performed with the XBLAST program, score = 50, wordlength = 3 to obtain amino acid sequences homologous to SMP protein molecules of the invention. To obtain gapped alignments for comparison purposes, Gapped BLAST can be utilized as described in Altschul *et al.*, (1997) *Nucleic Acids Res.* 25(17):3389-3402. When utilizing BLAST and Gapped BLAST programs, one of ordinary skill in the art will know how to optimize the parameters of the program (*e.g.*, XBLAST and NBLAST) for the specific sequence being analyzed.

Another example of a mathematical algorithm utilized for the comparison of sequences is the algorithm of Meyers and Miller ((1988) Comput. Appl. Biosci. 4: 11-17). Such an algorithm is incorporated into the ALIGN program (version 2.0) which is part of the GCG sequence alignment software package. When utilizing the ALIGN program for comparing amino acid sequences, a PAM120 weight residue table, a gap length penalty of 12, and a gap penalty of 4 can be used. Additional algorithms for sequence analysis are known in the art, and include ADVANCE and ADAM. described in Torelli and Robotti (1994) Comput. Appl. Biosci. 10:3-5; and FASTA, described in Pearson and Lipman (1988) P.N.A.S. 85:2444-8.

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The percent homology between two amino acid sequences can also be accomplished using the GAP program in the GCG software package (available at http://www.gcg.com), using either a Blosum 62 matrix or a PAM250 matrix, and a gap weight of 12, 10, 8, 6, or 4 and a length weight of 2, 3, or 4. The percent homology between two nucleic acid sequences can be accomplished using the GAP program in the GCG software package, using standard parameters, such as a gap weight of 50 and a length weight of 3.

A comparative analysis of the gene sequences of the invention with those present in Genbank has been performed using techniques known in the art (see, e.g., Bexevanis and Ouellette, eds. (1998) Bioinformatics: A Practical Guide to the Analysis of Genes and Proteins. John Wiley and Sons: New York). The gene sequences of the invention

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were compared to genes present in Genbank in a three-step process. In a first step, a BLASTN analysis (e.g., a local alignment analysis) was performed for each of the sequences of the invention against the nucleotide sequences present in Genbank, and the top 500 hits were retained for further analysis. A subsequent FASTA search (e.g., a combined local and global alignment analysis, in which limited regions of the sequences are aligned) was performed on these 500 hits. Each gene sequence of the invention was subsequently globally aligned to each of the top three FASTA hits, using the GAP program in the GCG software package (using standard parameters). In order to obtain correct results, the length of the sequences extracted from Genbank were adjusted to the length of the query sequences by methods well-known in the art. The results of this analysis are set forth in Table 4. The resulting data is identical to that which would have been obtained had a GAP (global) analysis alone been performed on each of the genes of the invention in comparison with each of the references in Genbank, but required significantly reduced computational time as compared to such a database-wide GAP (global) analysis. Sequences of the invention for which no alignments above the cutoff values were obtained are indicated on Table 4 by the absence of alignment information. It will further be understood by one of ordinary skill in the art that the GAP alignment homology percentages set forth in Table 4 under the heading "% homology (GAP)" are listed in the European numerical format, wherein a ',' represents a decimal point. For example, a value of "40,345" in this column represents "40.345%".

Example 12: Construction and Operation of DNA Microarrays

The sequences of the invention may additionally be used in the construction and application of DNA microarrays (the design, methodology, and uses of DNA arrays are well known in the art, and are described, for example, in Schena, M. et al. (1995) Science 270: 467-470; Wodicka, L. et al. (1997) Nature Biotechnology 15: 1359-1367; DeSaizieu, A. et al. (1998) Nature Biotechnology 16: 45-48; and DeRisi, J.L. et al. (1997) Science 278: 680-686).

DNA microarrays are solid or flexible supports consisting of nitrocellulose,
 nylon, glass, silicone, or other materials. Nucleic acid molecules may be attached to the surface in an ordered manner. After appropriate labeling, other nucleic acids or nucleic acid mixtures can be hybridized to the immobilized nucleic acid molecules, and the label

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may be used to monitor and measure the individual signal intensities of the hybridized molecules at defined regions. This methodology allows the simultaneous quantification of the relative or absolute amount of all or selected nucleic acids in the applied nucleic acid sample or mixture. DNA microarrays, therefore, permit an analysis of the expression of multiple (as many as 6800 or more) nucleic acids in parallel (see, e.g., Schena, M. (1996) *BioEssays* 18(5): 427-431).

The sequences of the invention may be used to design oligonucleotide primers which are able to amplify defined regions of one or more *C. glutamicum* genes by a nucleic acid amplification reaction such as the polymerase chain reaction. The choice and design of the 5' or 3' oligonucleotide primers or of appropriate linkers allows the covalent attachment of the resulting PCR products to the surface of a support medium described above (and also described, for example, Schena, M. *et al.* (1995) *Science* 270: 467-470).

Nucleic acid microarrays may also be constructed by *in situ* oligonucleotide synthesis as described by Wodicka, L. *et al.* (1997) *Nature Biotechnology* 15: 1359-1367. By photolithographic methods, precisely defined regions of the matrix are exposed to light. Protective groups which are photolabile are thereby activated and undergo nucleotide addition, whereas regions that are masked from light do not undergo any modification. Subsequent cycles of protection and light activation permit the synthesis of different oligonucleotides at defined positions. Small, defined regions of the genes of the invention may be synthesized on microarrays by solid phase oligonucleotide synthesis.

The nucleic acid molecules of the invention present in a sample or mixture of nucleotides may be hybridized to the microarrays. These nucleic acid molecules can be labeled according to standard methods. In brief, nucleic acid molecules (e.g., mRNA molecules or DNA molecules) are labeled by the incorporation of isotopically or fluorescently labeled nucleotides, e.g., during reverse transcription or DNA synthesis. Hybridization of labeled nucleic acids to microarrays is described (e.g., in Schena, M. et al. (1995) supra; Wodicka, L. et al. (1997), supra; and DeSaizieu A. et al. (1998), supra). The detection and quantification of the hybridized molecule are tailored to the specific incorporated label. Radioactive labels can be detected, for example, as

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described in Schena, M. et al. (1995) supra) and fluorescent labels may be detected, for example, by the method of Shalon et al. (1996) Genome Research 6: 639-645).

The application of the sequences of the invention to DNA microarray technology, as described above, permits comparative analyses of different strains of *C. glutamicum* or other Corynebacteria. For example, studies of inter-strain variations based on individual transcript profiles and the identification of genes that are important for specific and/or desired strain properties such as pathogenicity, productivity and stress tolerance are facilitated by nucleic acid array methodologies. Also, comparisons of the profile of expression of genes of the invention during the course of a fermentation reaction are possible using nucleic acid array technology.

Example 13: Analysis of the Dynamics of Cellular Protein Populations (Proteomics)

The genes, compositions, and methods of the invention may be applied to study the interactions and dynamics of populations of proteins, termed 'proteomics'. Protein populations of interest include, but are not limited to, the total protein population of *C. glutamicum* (e.g., in comparison with the protein populations of other organisms), those proteins which are active under specific environmental or metabolic conditions (e.g., during fermentation, at high or low temperature, or at high or low pH), or those proteins which are active during specific phases of growth and development.

Protein populations can be analyzed by various well-known techniques, such as gel electrophoresis. Cellular proteins may be obtained, for example, by lysis or extraction, and may be separated from one another using a variety of electrophoretic techniques. Sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) separates proteins largely on the basis of their molecular weight. Isoelectric focusing polyacrylamide gel electrophoresis (IEF-PAGE) separates proteins by their isoelectric point (which reflects not only the amino acid sequence but also posttranslational modifications of the protein). Another, more preferred method of protein analysis is the consecutive combination of both IEF-PAGE and SDS-PAGE, known as 2-D-gel electrophoresis (described, for example, in Hermann *et al.* (1998) *Electrophoresis* 19: 3217-3221; Fountoulakis *et al.* (1998) *Electrophoresis* 19: 1193-1202; Langen *et al.* (1997) *Electrophoresis* 18: 1184-1192; Antelmann *et al.* (1997) *Electrophoresis* 18:

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1451-1463). Other separation techniques may also be utilized for protein separation, such as capillary gel electrophoresis; such techniques are well known in the art.

Proteins separated by these methodologies can be visualized by standard techniques, such as by staining or labeling. Suitable stains are known in the art, and include Coomassie Brilliant Blue, silver stain, or fluorescent dyes such as Sypro Ruby (Molecular Probes). The inclusion of radioactively labeled amino acids or other protein precursors (e.g., ³⁵S-methionine, ³⁵S-cysteine, ¹⁴C-labelled amino acids, ¹⁵N-amino acids, ¹⁵NO₃ or ¹⁵NH₄⁺ or ¹³C-labelled amino acids) in the medium of *C. glutamicum* permits the labeling of proteins from these cells prior to their separation. Similarly, fluorescent labels may be employed. These labeled proteins can be extracted, isolated and separated according to the previously described techniques.

Proteins visualized by these techniques can be further analyzed by measuring the amount of dye or label used. The amount of a given protein can be determined quantitatively using, for example, optical methods and can be compared to the amount of other proteins in the same gel or in other gels. Comparisons of proteins on gels can be made, for example, by optical comparison, by spectroscopy, by image scanning and analysis of gels, or through the use of photographic films and screens. Such techniques are well-known in the art.

To determine the identity of any given protein, direct sequencing or other standard techniques may be employed. For example, N- and/or C-terminal amino acid sequencing (such as Edman degradation) may be used, as may mass spectrometry (in particular MALDI or ESI techniques (see, e.g., Langen et al. (1997) Electrophoresis 18: 1184-1192)). The protein sequences provided herein can be used for the identification of C. glutamicum proteins by these techniques.

The information obtained by these methods can be used to compare patterns of protein presence, activity, or modification between different samples from various biological conditions (e.g., different organisms, time points of fermentation, media conditions, or different biotopes, among others). Data obtained from such experiments alone, or in combination with other techniques, can be used for various applications, such as to compare the behavior of various organisms in a given (e.g., metabolic) situation, to increase the productivity of strains which produce fine chemicals or to increase the efficiency of the production of fine chemicals.

- 133 -

Equivalents

Those of ordinary skill in the art will recognize, or will be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the following claims.

What is claimed:

- An isolated nucleic acid molecule from Corynebacterium glutamicum encoding an
 SMP protein, or a portion thereof, provided that the nucleic acid molecule does not consist of any of the F-designated genes set forth in Table 1.
 - 2. The isolated nucleic acid molecule of claim 1, wherein said nucleic acid molecule encodes an SMP protein involved in the production of a fine chemical.

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3. An isolated *Corynebacterium glutamicum* nucleic acid molecule selected from the group consisting of those sequences set forth as odd-numbered SEQ ID NOs of the Sequence Listing, or a portion thereof, provided that the nucleic acid molecule does not consist of any of the F-designated genes set forth in Table 1.

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4. An isolated nucleic acid molecule which encodes a polypeptide sequence selected from the group consisting of those sequences set forth as even-numbered SEQ ID NOs of the Sequence Listing,, provided that the nucleic acid molecule does not consist of any of the F-designated genes set forth in Table 1.

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- 5. An isolated nucleic acid molecule which encodes a naturally occurring allelic variant of a polypeptide selected from the group of amino acid sequences consisting of those sequences set forth in as even-numbered SEQ ID NOs of the Sequence Listing, provided that the nucleic acid molecule does not consist of any of the F-designated genes set forth in Table 1.
- 6. An isolated nucleic acid molecule comprising a nucleotide sequence which is at least 50% homologous to a nucleotide sequence selected from the group consisting of those sequences set forth as odd-numbered SEQ ID NOs of the Sequence Listing, or a portion thereof, provided that the nucleic acid molecule does not consist of any of the F-designated genes set forth in Table 1.

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- 7. An isolated nucleic acid molecule comprising a fragment of at least 15 nucleotides of a nucleic acid comprising a nucleotide sequence selected from the group consisting of those sequences set forth as odd-numbered SEQ ID NOs of the Sequence Listing, provided that the nucleic acid molecule does not consist of any of the F-designated genes set forth in Table 1.
- 8. An isolated nucleic acid molecule which hybridizes to the nucleic acid molecule of any one of claims 1-7 under stringent conditions.
- 9. An isolated nucleic acid molecule comprising the nucleic acid molecule of any one of claims 1-8 or a portion thereof and a nucleotide sequence encoding a heterologous polypeptide.
 - 10. A vector comprising the nucleic acid molecule of any one of claims 1-9.
 - 11. The vector of claim 10, which is an expression vector.
 - 12. A host cell transfected with the expression vector of claim 11.
- 20 13. The host cell of claim 12, wherein said cell is a microorganism.
 - 14. The host cell of claim 13, wherein said cell belongs to the genus *Corynebacterium* or *Brevibacterium*.
- 25 15. The host cell of claim 12, wherein the expression of said nucleic acid molecule results in the modulation in production of a fine chemical from said cell.
- 16. The host cell of claim 15, wherein said fine chemical is selected from the group consisting of: organic acids, proteinogenic and nonproteinogenic amino acids, purine
 and pyrimidine bases, nucleosides, nucleotides, lipids, saturated and unsaturated fatty acids, diols, carbohydrates, aromatic compounds, vitamins, cofactors, polyketides, and enzymes.

- 17. A method of producing a polypeptide comprising culturing the host cell of claim 12 in an appropriate culture medium to, thereby, produce the polypeptide.
- 5 18. An isolated SMP polypeptide from *Corynebacterium glutamicum*, or a portion thereof.
 - 19. The polypeptide of claim 18, wherein said polypeptide is involved in the production of a fine chemical.

20. An isolated polypeptide comprising an amino acid sequence selected from the group consisting of those sequences set forth as even-numbered SEQ ID NOs of the Sequence Listing, provided that the amino acid sequence is not encoded by any of

the F-designated genes set forth in Table 1,

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21. An isolated polypeptide comprising a naturally occurring allelic variant of a polypeptide comprising an amino acid sequence selected from the group consisting of those sequences set forth as even-numbered SEQ ID NOs of the Sequence Listing, or a portion thereof, provided that the amino acid sequence is not encoded by any of the F-designated genes set forth in Table 1.

- 22. The isolated polypeptide of any of claims 18-21, further comprising heterologous amino acid sequences.
- 25 23. An isolated polypeptide which is encoded by a nucleic acid molecule comprising a nucleotide sequence which is at least 50% homologous to a nucleic acid selected from the group consisting of those sequences set forth as odd-numbered SEQ ID NOs of the Sequence Listing, provided that the nucleic acid molecule does not consist of any of the F-designated nucleic acid molecules set forth in Table 1.
 - 24. An isolated polypeptide comprising an amino acid sequence which is at least 50% homologous to an amino acid sequence selected from the group consisting of those

sequences as even-numbered SEQ ID NOs of the Sequence Listing, provided that the amino acid sequence is not encoded by any of the F-designated genes set forth in Table 1.

- 5 25. A method for producing a fine chemical, comprising culturing a cell containing a vector of claim 12 such that the fine chemical is produced.
 - 26. The method of claim 25, wherein said method further comprises the step of recovering the fine chemical from said culture.

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- 27. The method of claim 25, wherein said method further comprises the step of transfecting said cell with the vector of claim 11 to result in a cell containing said vector.
- 28. The method of claim 25, wherein said cell belongs to the genus *Corynebacterium* or *Brevibacterium*.
 - 29. The method of claim 25, wherein said cell is selected from the group consisting of: Corynebacterium glutamicum, Corynebacterium herculis, Corynebacterium, lilium,
- Corynebacterium acetoacidophilum, Corynebacterium acetoglutamicum,
 Corynebacterium acetophilum, Corynebacterium ammoniagenes, Corynebacterium
 fujiokense, Corynebacterium nitrilophilus, Brevibacterium ammoniagenes,
 Brevibacterium butanicum, Brevibacterium divaricatum, Brevibacterium flavum,
 Brevibacterium healii, Brevibacterium ketoglutamicum, Brevibacterium
- 25 ketosoreductum, Brevibacterium lactofermentum, Brevibacterium linens, Brevibacterium paraffinolyticum, and those strains set forth in Table 3.
 - 30. The method of claim 25, wherein expression of the nucleic acid molecule from said vector results in modulation of production of said fine chemical.
 - 31. The method of claim 25, wherein said fine chemical is selected from the group consisting of: organic acids, proteinogenic and nonproteinogenic amino acids, purine

and pyrimidine bases, nucleosides, nucleotides, lipids, saturated and unsaturated fatty acids, diols, carbohydrates, aromatic compounds, vitamins, cofactors, polyketides, and enzymes.

- 5 32. The method of claim 25, wherein said fine chemical is an amino acid.
 - 33. The method of claim 32, wherein said amino acid is drawn from the group consisting of: lysine, glutamate, glutamine, alanine, aspartate, glycine, serine, threonine, methionine, cysteine, valine, leucine, isoleucine, arginine, proline, histidine, tyrosine, phenylalanine, and tryptophan.
 - 34. A method for producing a fine chemical, comprising culturing a cell whose genomic DNA has been altered by the inclusion of a nucleic acid molecule of any one of claims 1-9.

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- 35. A method for diagnosing the presence or activity of *Corynebacterium* diphtheriae in a subject, comprising detecting the presence of one or more of SEQ ID NOs 1 through 782 of the Sequence Listing in the subject, provided that the sequences are not or are not encoded by any of the F-designated sequences set forth in Table 1, thereby diagnosing the presence or activity of *Corynebacterium diphtheriae* in the subject.
- 36. A host cell comprising a nucleic acid molecule selected from the group consisting of the nucleic acid molecules set forth as odd-numbered SEQ ID NOs of the
 25 Sequence Listing, wherein the nucleic acid molecule is disrupted.
 - 35. 37. A host cell comprising a nucleic acid molecule selected from the group consisting of the nucleic acid molecules set forth as odd-numbered SEQ ID NOs in the Sequence Listing, wherein the nucleic acid molecule comprises one or more nucleic acid modifications from the sequence set forth as odd-numbered SEQ ID NOs of the Sequence Listing s.

38. A host cell comprising a nucleic acid molecule selected from the group consisting of the nucleic acid molecules set forth as odd-numbered SEQ ID NOs of the Sequence Listing, wherein the regulatory region of the nucleic acid molecule is modified relative to the wild-type regulatory region of the molecule.

SEQUENCE LISTING

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													tcc Ser			163
													gca Ala 35			211
													aag Lys			259
													ttc Phe			307
													gag Glu			355
													gcc Ala			403
													gcc Ala 115			451
													gat Asp			499
eta c	ate	gge	atσ	aat	aac	gaa	aac	cat	atc	aac	tee	cta	ttc	cct	cac	547

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gct Ala	gct Ala 215	gga Gly	gct Ala	acc Thr	gga Gly	tct Ser 220	gag Glu	gaa Glu	acg Thr	gta Val	ttg Leu 225	ttc Phe	ttg Leu	gct Ala	gat Asp	787
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					cac His											403
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1	C.c.	7	+	5	63	63	**- *	.	10	17- 7	D	7	n 7 -	15	m	
rne	ser	Arg	Leu 20	GTÀ	Glu	GIn	val	Leu 25	Ата	val	Pro	Asp	A1a 30	Asp	Trp	

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25 30 35

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atc Ile	cct Pro	aac Asn	tac Tyr	ctc Leu 490	Gly	cca Pro	ttg Leu	ctt Leu	ggc Gly 495	tcc Ser	gag Glu	cgt Arg	ctg Leu	tca Ser 500	Glu	1603
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						cct Pro 140										547
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						ctg Leu 220									acc Thr	787
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						aag Lys										883
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						acc Thr 300										1027
_	_					atc Ile					-	•				1075
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Val Asn His Glu Thr Ala Glu Leu Ala Leu Asp His Ala Ala Cys Ile 150 Gly Cys Gly Ala Cys Val Ala Ala Cys Pro Asn Gly Ala Ala His Leu 170 Phe Thr Gly Ala Lys Leu Val His Leu Ser Leu Leu Pro Leu Gly Lys 185 Glu Glu Arg Gly Leu Arg Ala Arg Lys Met Val Asp Glu Met Glu Thr 200 Asn Phe Gly His Cys Ser Leu Tyr Gly Glu Cys Ala Asp Val Cys Pro 215 Ala Gly Ile Pro Leu Thr Ala Val Ala Ala Val Thr Lys Glu Arg Ala 235 225 230 · Arg Ala Ala Phe Arg Gly Lys Asp Asp 245 <210> 17 <211> 1530 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1507) <223> RXA01535 <400> 17 acceaectea etetaggggt ggaeteeagt gtttegegae aacacaatga gtaagettgt 60 qacaqccqta tttaattctc agtaagaaat gagtgatttc atg acc gag cag gaa Met Thr Glu Gln Glu 1 ttc cgt att gag cac gac acc atg ggt gaa gtg aag gtt cca gca aag Phe Arg Ile Glu His Asp Thr Met Gly Glu Val Lys Val Pro Ala Lys 10 211 get etg tag cag gea cag ace cag ege get gtt gag aac tte eet ate Ala Leu Trp Gln Ala Gln Thr Gln Arg Ala Val Glu Asn Phe Pro Ile 25 tot ggt cgt ggt ctg gaa too goa cag ato cgc gca atg ggt ctg ctg 259 Ser Gly Arg Gly Leu Glu Ser Ala Gln Ile Arg Ala Met Gly Leu Leu 40 45 307 aag gca gct tgt gcg cag gta aac aag gac tcc ggt gcg ctg gat gca Lys Ala Ala Cys Ala Gln Val Asn Lys Asp Ser Gly Ala Leu Asp Ala 55 gag aag gca gat gcc atc att gca gct ggt aag gag atc gcg tcc ggt 355 Glu Lys Ala Asp Ala Ile Ile Ala Ala Gly Lys Glu Ile Ala Ser Gly 80 75 aag cat gac gct gag ttc cca att gat gtg ttc cag act ggt tcc ggt 403 Lys His Asp Ala Glu Phe Pro Ile Asp Val Phe Gln Thr Gly Ser Gly

Ala 150	Glu	Leu	Ala	Leu	Asp 155	His	Ala	Ala	Cys	Ile 160	Gly	Cys	Gly	Ala	Cys 165	
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	gtt Val															691
	gca Ala					Asp										739
	ctc Leu 215															787
	gct Ala		-	-	-		-	-	_		_	-				835
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Lys Glu Pro Phe Ala Phe Ala Ser Asp Cys Arg Glu Gly Ile Cys Gly 50 55 60

Thr Cys Gly Leu Leu Val Asn Gly Arg Pro His Gly Ala Asp Gln Asn 65 70 75 80

Lys Pro Ala Cys Ala Gln Arg Leu Val Ser Tyr Lys Glu Gly Asp Thr 85 90 95

Leu Lys Ile Glu Pro Leu Arg Ser Ala Ala Tyr Pro Val Ile Lys Asp 100 105 110

Met Val Val Asp Arg Ser Ala Leu Asp Arg Val Met Glu Gln Gly Gly 115 120 125

Tyr Val Thr Ile Asn Ala Gly Thr Ala Pro Asp Ala Asp Thr Leu His 130 135 140

21

475

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Glu 390	atc Ile				_	_	_		_						-	1315
	gcc Ala															1363
	ttc Phe															1411
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	cct Pro 455															1507
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_	cct Pro		_		tago	catc	tgc (cccti	taca	aa at	c					1593
	0> 14															
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145

547

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.

140

135

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WO 01/00844	PCT/IB00/00943

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Thr Trp Gln Ser Lys Thr Ile Leu Met Ser Glu Ser Leu Arg Asn Asp 305 310 315 320

Gly Arg Ile Trp Ser Pro Lys Glu Pro Asn Asp Asn Arg Asp Pro Asn 325 330 335

Thr Ile Pro Glu Asp Glu Arg Asp Tyr Phe Leu Glu Arg Arg Tyr Pro 340 345 350

Ala Phe Gly Asn Leu Val Pro Arg Asp Val Ala Ser Arg Ala Ile Ser 355 360 365

Gln Gln Ile Asn Ala Gly Leu Gly Val Gly Pro Leu Asn Asn Ala Ala 370 375 380

Tyr Leu Asp Phe Arg Asp Ala Thr Glu Arg Leu Gly Gln Asp Thr Ile 385 390 395 400

Arg Glu Arg Tyr Ser Asn Leu Phe Thr Met Tyr Glu Glu Ala Ile Gly
405 410 415

Glu Asp Pro Tyr Ser Ser Pro Met Arg Ile Ala Pro Thr Cys His Phe 420 425 430

Thr Met Gly Gly Leu Trp Thr Asp Phe Asn Glu Met Thr Ser Leu Pro 435 440 445

Gly Leu Phe Cys Ala Gly Glu Ala Ser Trp Thr Tyr His Gly Ala Asn 450 455 460

Arg Leu Gly Ala Asn Ser Leu Leu Ser Ala Ser Val Asp Gly Trp Phe 465 470 480

Thr Leu Pro Phe Thr Ile Pro Asn Tyr Leu Gly Pro Leu Leu Gly Ser 485 490 495

Glu Arg Leu Ser Glu Asp Ala Pro Glu Ala Gln Ala Ala Ile Ala Arg 500 505 510

Ala Gln Ala Arg Ile Asp Arg Leu Met Gly Asn Arg Pro Glu Trp Val 515 520 525

Gly Asp Asn Val His Gly Pro Glu Tyr Tyr His Arg Gln Leu Gly Asp 530 540

Ile Leu Tyr Phe Ser Cys Gly Val Ser Arg Asn Val Glu Asp Leu Gln 545 550 555 560

Asp Gly Ile Asn Lys Ile Arg Ala Leu Arg Asp Asp Phe Trp Lys Asn 565 570 575

Met Arg Ile Thr Gly Ser Thr Asp Glu Met Asn Gln Val Leu Glu Tyr 580 585 590

Ala Ala Arg Val Ala Asp Tyr Ile Asp Leu Gly Glu Leu Met Cys Val 595 600 605

Asp Ala Leu Asp Arg Asp Glu Ser Cys Gly Ala His Phe Arg Asp Asp 610 620

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<213> Corynebacterium glutamicum

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Pro Ala Gly Val Pro Thr Lys Asp Met Trp Glu Tyr Gln Lys Asp His 35 40 45

Met Asn Leu Val Ser Pro Leu Asn Arg Arg Lys Phe Arg Val Leu Val 50 55 60

Val Gly Thr Gly Leu Ser Gly Gly Ala Ala Ala Ala Ala Leu Gly Glu 65 70 75 80

Leu Gly Tyr Asp Val Lys Ala Phe Thr Tyr His Asp Ala Pro Arg Arg 85 90 95

Ala His Ser Ile Ala Ala Gln Gly Gly Val Asn Ser Ala Arg Gly Lys 100 105 110

Lys Val Asp Asn Asp Gly Ala Tyr Arg His Val Lys Asp Thr Val Lys 115 120 125

Gly Gly Asp Tyr Arg Gly Arg Glu Ser Asp Cys Trp Arg Leu Ala Val 130 140

Glu Ser Val Arg Val Ile Asp His Met Asn Ala Ile Gly Ala Pro Phe 145 150 155 160

Ala Arg Glu Tyr Gly Gly Ala Leu Ala Thr Arg Ser Phe Gly Gly Val 165 170 175

Gln Val Ser Arg Thr Tyr Tyr Thr Arg Gly Gln Thr Gly Gln Gln Leu 180 185 190

Gln Phe Ser Thr Ala Ser Ala Leu Gln Arg Gln Ile His Leu Gly Ser 195 200 205

Val Glu Ile Phe Thr His Asn Glu Met Val Asp Val Ile Val Thr Glu 210 215 220

Arg Asn Gly Glu Lys Arg Cys Glu Gly Leu Ile Met Arg Asn Leu Ile 225 230 235 240

Thr Gly Glu Leu Thr Ala His Thr Gly His Ala Val Ile Leu Ala Thr 245 250 255

Gly Gly Tyr Gly Asn Val Tyr His Met Ser Thr Leu Ala Lys Asn Ser

Asn Ala Ser Ala Ile Met Arg Ala Tyr Glu Ala Gly Ala Tyr Phe Ala 275 280 285

Ser Pro Ser Phe Ile Gln Phe His Pro Thr Gly Leu Pro Val Asn Ser 290 295 300

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Phe Ser Gly Ala Met Arg Val Ile Ala Val Ser Leu Tyr Lys Ile Ala 275 280

Asn Asp Ile Arg Leu Met Gly Ser Gly Pro Leu Thr Gly Leu Gly Glu 295

Ile Arg Leu Pro Asp Leu Gln Pro Gly Ser Ser Ile Met Pro Gly Lys

Val Asn Pro Val Leu Cys Glu Thr Ala Thr Gln Val Ser Ala Gln Val 330 325

Ile Gly Asn Asp Ala Ala Val Ala Phe Ser Gly Thr Gln Gly Gln Phe 340

Glu Leu Asn Val Phe Ile Pro Val Met Ala Arg Asn Val Leu Glu Ser 355 360

Ala Arg Leu Leu Ala Asn Thr Ser Arg Val Phe Ala Thr Arg Leu Val 370 375

Asp Gly Ile Glu Pro Asn Glu Ala His Met Lys Glu Leu Ala Glu Ser

Ser Pro Ser Ile Val Thr Pro Leu Asn Ser Ala Ile Gly Tyr Glu Ala 410

Ala Ala Lys Val Ala Lys Thr Ala Leu Ala Glu Gly Lys Thr Ile Arg 420

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						gct Ala										643
						tac Tyr										691
						acg Thr										739
						gtc Val 220										787
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						gta Val									Pro	883
						ttc Phe										931
						gac Asp										979
						caa Gln 300										1027
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Gly Asp Ser Leu Val Gln Ala Asp Leu Trp Gly His Pro Ser His Gly 35 40 45

Val Leu Arg Leu Pro Trp Tyr Val Arg Arg Leu His Ser Gly Ala Met 50 55 60

Thr Thr His Ala His Val Glu Val Leu Asn Asp Leu Gly Ala Val Leu 65 70 75 80

Ala Leu Asp Gly His Asn Gly Ile Gly Gln Val Leu Ala Asp His Ala 85 90 95

Arg Lys Glu Ala Val Thr Arg Ala Met Met Phe Gly Ile Gly Ala Val 100 105 110

Ser Val Arg Asn Ser Asn His Phe Gly Thr Ala Met Tyr Tyr Thr Arg 115 120 125

Lys Ala Ala Ala Gln Gly Cys Val Ser Ile Leu Thr Thr Asn Ala Ser 130 135 140

Pro Ala Met Ala Pro Trp Gly Gly Arg Glu Lys Arg Ile Gly Thr Asn 145 150 155 160

Pro Trp Ser Ile Ala Ala Pro Phe Gly Glu Thr Ala Thr Val Val Asp 165 170 175

Ile Ala Asn Thr Ala Val Ala Arg Gly Lys Ile Tyr His Ala Arg Gln 180 185 190

Thr Asn Met Pro Ile Pro Glu Thr Trp Ala Ile Thr Ser Glu Gly Ala 195 200 205

Pro Thr Thr Asp Pro Ala Glu Ala Ile Asn Gly Val Val Leu Pro Met 210 215 220

Ala Gly His Lys Gly Tyr Ala Ile Ser Phe Met Met Asp Val Leu Ser 225 230 235 240

Gly Val Leu Thr Gly Ser Gln His Ser Thr Lys Val His Gly Pro Tyr 245 250 255

Asp Pro Thr Pro Pro Gly Gly Ala Gly His Leu Phe Ile Ala Leu Asp 260 265 270

Val Ala Ala Phe Arg Asp Pro Gln Asp Phe Asp Asp Ala Leu Ser Asp 275 280 285

Leu Val Gly Glu Val Lys Ser Thr Pro Lys Ala Gln Asn Thr Glu Glu 290 295 300

Ile Phe Tyr Pro Gly Glu Ser Glu Asp Arg Ala His Arg Lys Asn Ser 305 310 315 320

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Leu Ala Ile Glu Asn His Val Val Thr His Arg 340 345

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Gly Ala Glu Gly Val Ala Met Glu Leu Leu Asp Ser Ala Phe Pro Leu
55 60 65

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gag cgc gca gat ttg ctg gct aac aac ggc aag att ttc gga cct caa 451 Glu Arg Ala Asp Leu Leu Ala Asn Asn Gly Lys Ile Phe Gly Pro Gln 105 110 115

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cca gat gtt cca gca tcc cgc ttc aac gca atg atg cgc ctt gat cac 595

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gaa Glu	ttt Phe	aac Asn	aac Asn 185	att Ile	gtg Val	gtc Val	tgg Trp	gga Gly 190	aat Asn	cac His	tcc Ser	gca Ala	acc Thr 195	cag Gln	ttc Phe	691
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gtt Val	gat Asp 215	cac His	gat Asp	tgg Trp	tat Tyr	gtg Val 220	gag Glu	gag Glu	ttc Phe	att Ile	cct Pro 225	cgc Arg	gtg Val	gct Ala	aac Asn	787
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gag Glu	gcg Ala	tgg Trp	tcc Ser 265	tct Ser	gcg Ala	gca Ala	att Ile	cct Pro 270	tcc Ser	acc Thr	ggt Gly	gca Ala	tac Tyr 275	ggc Gly	att Ile	931
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35 40 45

Pro Gln Ala Leu Gly Gly Ala Glu Gly Val Ala Met Glu Leu Leu Asp 50 55 60

Ser Ala Phe Pro Leu Leu Arg Asn Ile Thr Ile Thr Ala Asp Ala Asn 65 70 75 80

Glu Ala Phe Asp Gly Ala Asn Ala Ala Phe Leu Val Gly Ala Lys Pro 85 90 95

Arg Gly Lys Gly Glu Glu Arg Ala Asp Leu Leu Ala Asn Asn Gly Lys 100 105 110

Ile Phe Gly Pro Gln Gly Lys Ala Ile Asn Asp Asn Ala Ala Asp Asp 115 120 125

Ile Arg Val Leu Val Val Gly Asn Pro Ala Asn Thr Asn Ala Leu Ile 130 135 140

Ala Ser Ala Ala Ala Pro Asp Val Pro Ala Ser Arg Phe Asn Ala Met 145 150 155 160

Met Arg Leu Asp His Asn Arg Ala Ile Ser Gln Leu Ala Thr Lys Leu 165 170 175

Gly Arg Gly Ser Ala Glu Phe Asn Asn Ile Val Val Trp Gly Asn His 180 185 190

Ser Ala Thr Gln Phe Pro Asp Ile Thr Tyr Ala Thr Val Gly Glu 195 200 205

Lys Val Thr Asp Leu Val Asp His Asp Trp Tyr Val Glu Glu Phe Ile 210 215 220

Pro Arg Val Ala Asn Arg Gly Ala Glu Ile Ile Glu Val Arg Gly Lys 225 230 235 240

Ser Ser Ala Ala Ser Ala Ala Ser Ser Ala Ile Asp His Met Arg Asp 245 250 255

Trp Val Gln Gly Thr Glu Ala Trp Ser Ser Ala Ala Ile Pro Ser Thr 260 265 270

Gly Ala Tyr Gly Ile Pro Glu Gly Ile Phe Val Gly Leu Pro Thr Val 275 280 285

Ser Arg Asn Gly Glu Trp Glu Ile Val Glu Gly Leu Glu Ile Ser Asp 290 295 300

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<211> 1092

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Gly	ctt Leu	gto Val	gac Asp 25	Glu	tco Ser	ggg Gly	cgc Arg	ato Ile 30	Val	acc Thr	agt Ser	ttg Leu	tcg Ser 35	Ala	ccg	211
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gcg Ala 70	gga Gly	ttt Phe	ttg Leu	gat Asp	cct Pro 75	Glu	tgc Cys	gag Glu	gtt Val	gtt Val 80	Arg	ttt Phe	gcc Ala	ccg Pro	cac His 85	355
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gag Glu	cat His	cgt Arg 120	ttt Phe	ggt Gly	gca Ala	gct Ala	caa Gln 125	ggc Gly	gct Ala	gac Asp	aac Asn	tgg Trp 130	gtt Val	ttg Leu	ttg Leu	499
gca Ala	ctc Leu 135	ggc Gly	act Thr	gga Gly	att Ile	ggt Gly 140	gca Ala	gcg Ala	ctg Leu	att Ile	gaa Glu 145	aaa Lys	ggc Gly	gaa Glu	att Ile	547
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egt Arg	tac Tyr	tgt Cys	tcc Ser 185	ggt Gly	act Thr	gcc Ala	ttg Leu	gtt Val 190	tac Tyr	act Thr	gcg Ala	cgt Arg	gaa Glu 195	ttg Leu	gct Ala	691

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						gga Gly 220										787
						gcc Ala										835
						att Ile										883
						tcc Ser										931
						acc Thr										979
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Ser Ala Ala Trp Gly Glu His Arg Phe Gly Ala Ala Gln Gly Ala Asp Asn Trp Val Leu Leu Ala Leu Gly Thr Gly Ile Gly Ala Ala Leu Ile 135 Glu Lys Gly Glu Ile Tyr Arg Gly Ala Tyr Gly Thr Ala Pro Glu Phe Gly His Leu Arg Val Val Arg Gly Gly Arg Ala Cys Ala Cys Gly Lys 170 Glu Gly Cys Leu Glu Arg Tyr Cys Ser Gly Thr Ala Leu Val Tyr Thr 180 185 Ala Arg Glu Leu Ala Ser His Gly Ser Phe Arg Asn Ser Gly Leu Phe 200 Asp Lys Ile Lys Ala Asp Pro Asn Ser Ile Asn Gly Lys Thr Ile Thr 215 Ala Ala Arg Gln Glu Asp Pro Leu Ala Leu Ala Val Leu Glu Asp Phe Ser Glu Trp Leu Gly Glu Thr Leu Ala Ile Ile Ala Asp Val Leu 250 Asp Pro Gly Met Ile Ile Gly Gly Gly Leu Ser Asn Ala Ala Asp 2.60 265 Leu Tyr Leu Asp Arg Ser Val Asn His Tyr Ser Thr Arg Ile Val Gly 280 Ala Gly Tyr Arg Pro Leu Ala Arg Val Ala Thr Ala Gln Leu Gly Ala 290 295 Asp Ala Gly Met Ile Gly Val Ala Asp Leu Ala Arg Arg Ser Val Val 310 315 Glu Ala Asn <210> 25 <211> 1785 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1762) <223> RXA01814 <400> 25 tgttaagcca ccctactccg tgaattttgc cgtatctcgt gcgcacaatt gcttttgaqq 60 ggaagatgaa gagaaagtat tggtgtttta aggagcaaac atg gca cat gaa cqc Met Ala His Glu Arg

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					acc Thr											211
					acc Thr											259
					cac His											307
					cca Pro 75											355
					ctg Leu											403
					gac Asp											451
					gca Ala											499
					acc Thr											547
					tcc Ser 155											595
					ggt Gly											643
					aac Asn											691
					tcc Ser											739
					tac Tyr											787
					gct Ala 235											835
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Phe Ala Leu Asp Ser Ala Phe Asn Glu Asp His Ile Leu Ala Thr Thr

Gln Ala Ile Val Asp Tyr Arg Asn Gln Gln Pro Lys Asn Trp Val Gly

Pro Leu Phe Ile Gly Arg Asp Thr His Ala Leu Ser Glu Pro Ala Met

Ile Ser Ala Leu Glu Val Leu Ile Ala Asn Asp Val Glu Val Leu Val 105

Asp Ala Asp Gly Arg Tyr Thr Pro Thr Pro Ala Val Ser His Ala Ile 115 120

Leu Arg His Asn Asp Gly Ile Ile Leu Gly Thr Ala Gly Pro Ser Arg

Pro Tyr Ala Asp Gly Ile Val Ile Thr Pro Ser His Asn Pro Pro Arg 145

Asp Gly Gly Phe Lys Tyr Asn Pro Ala Asn Gly Gly Pro Ala Asp Thr

Asp Ala Thr Asp Trp Ile Ala Asn Arg Ala Asn Asp Ile Leu Arg Gly 185 180

Asp Leu Ala Asp Val Lys Arg Val Pro Val Ser Gly Val Leu Asp Glu 195 200 205

Arg Thr Thr Ala Tyr Asp Phe Lys Gly Ile Tyr Ile Ala Asp Leu Pro 210 215 220

Asn Val Val Asn Ile Asp Ala Ile Arg Glu Ala Gly Val Arg Ile Gly 225 230 235 240

Ala Asp Pro Met Gly Gly Ala Ser Val Asp Tyr Trp Gly Ala Ile Ala 245 250 255

Glu Thr His Gly Leu Asn Leu Thr Val Val Asn Pro His Val Asp Ser 260 265 270

Thr Phe Arg Phe Met Thr Leu Asp Thr Asp Gly Lys Ile Arg Met Asp 275 280 285

Cys Ser Ser Pro His Ala Met Ala Ser Leu Ile Asp Asn Arg Asp Lys 290 295 300

Phe Asp Val Ala Thr Gly Asn Asp Ala Asp Ala Asp Arg His Gly Ile 305 310 315 320

Val Thr Pro Asp Ala Gly Leu Met Asn Pro Asn His Tyr Leu Ala Val 325 330 335

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Ala Val Gly Lys Thr Leu Val Ser Ser Ser Met Ile Asp Arg Val Val 355 360 365

Ala Gln Leu Gly Arg Thr Leu Val Glu Val Pro Val Gly Phe Lys Trp 370 375 380

Phe Val Pro Gly Leu Ile Ser Gly Glu Ile Gly Phe Gly Glu Glu 385 390 395 400

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Asp Lys Asp Gly Leu Ile Leu Asp Leu Leu Ala Ala Glu Ile Ile Ala 420 425 430

Val Thr Gly Lys Thr Pro Ser Gln Arg Tyr Ala Glu Leu Ala Glu Glu 435 440 445

Phe Gly Ala Pro Ala Tyr Ala Arg Thr Asp Ala Glu Ala Asn Arg Glu 450 455 460

Gln Lys Ala Ile Leu Lys Ala Leu Ser Pro Glu Gln Val Thr Ala Thr 465 470 475 480

Glu Leu Ala Gly Glu Ala Ile Thr Ala Lys Leu Thr Glu Ala Pro Gly
485 490 495

Asn Gly Ala Ala Ile Gly Gly Leu Lys Val Thr Thr Glu Asn Ala Trp 500 505 510

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155

Ala Gly Ala Thr Val Ile Ala Ile His Asn Lys Pro Asp Ser Tyr Asn

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150

145

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Gly 65	His	Lys	Leu	Pro	Asp 70	His	Val	Glu	Asp	Glu 75	Ile	Glu	Arg	Val	Met 80	
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						ttg Leu										211
						cgt Arg										259
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						cct Pro										451
						gat Asp										499
						ggc Gly 140										547
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						cgg Arg										739
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						ttt Phe										835
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Gly Ala Gly Phe Glu Val Thr Leu Leu Pro Thr Pro Ser Pro Thr Pro 100 105 110

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Ile Thr Ala Ser His Asn Gly Ala Ala Asp Asn Gly Tyr Lys Val Phe

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Ala His Ile Asn Ala Val Glu Asp Pro Ile Arg Val Pro Arg Val Thr 165 170 175

Val Arg Pro Thr Ala Asp Gln Leu Arg Arg Tyr Val Asp Glu Met Val 180 185 190

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Arg Ala Met Ala Asn Ala Phe Gln Phe Ala Gly Phe Pro His Thr His 225 230 235 240

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35

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Met 1 Tyr Asn Leu 65 Gly Leu Ala	Ala Ser Arg Ser 50 Thr Glu Arg	Asp Ala 35 Lys Glu His Leu Asp 115	Phe 20 Glu Asn Glu Leu Pro 100 Val	5 Gln Lys Leu Ser Asn 85 Ala	Ala Tyr Leu Gly 70 Asn Glu	Thr Asp 55 Leu Thr Ala	Thr Phe 40 Asp Arg Glu Asp Leu 120	Leu 25 Ser Ala Glu Asp Leu 105 Gly	10 Arg Ala Thr Arg Ser Arg	Glu Ala Leu Ile 75 Ala Val	Leu Gly Thr 60 Asp Val Asp	Phe Leu 45 Lys Ala Leu Gly Asp 125	Lys 30 His Leu Met His Gln 110	15 Glu Val Leu Phe Thr 95 Asp	Glu Asp Ala Ala 80 Ala Val	

Ala Thr Lys Ala Leu Arg Ala Tyr Ala Thr Ala Gly Ile Ser Ala Glu 165 170 175

- Phe Val Ser Asn Val Asp Pro Ala Asp Leu Val Ser Val Leu Glu Asp 180 185 190
- Leu Asp Ala Glu Ser Thr Leu Phe Val Ile Ala Ser Lys Thr Phe Thr 195 200 205
- Thr Gln Glu Thr Leu Ser Asn Ala Arg Ala Arg Ala Trp Leu Val 210 215 220
- Glu Lys Leu Gly Glu Glu Ala Val Ala Lys His Phe Val Ala Val Ser 225 230 235 240
- Thr Asn Ala Glu Lys Val Ala Glu Phe Gly Ile Asp Thr Asp Asn Met
 . 245 250 255
- Phe Gly Phe Trp Asp Trp Val Gly Gly Arg Tyr Ser Val Asp Ser Ala 260 265 270
- Val Gly Leu Ser Leu Met Ala Val Ile Gly Pro Arg Asp Phe Met Arg 275 280 285
- Phe Leu Gly Gly Phe His Ala Met Asp Glu His Phe Arg Thr Thr Lys 290 295 300
- Phe Glu Glu Asn Val Pro Ile Leu Met Ala Leu Leu Gly Val Trp Tyr 305 310 315 320
- Ser Asp Phe Tyr Gly Ala Glu Thr His Ala Val Leu Pro Tyr Ser Glu 325 330 335
- Asp Leu Ser Arg Phe Ala Ala Tyr Leu Gln Gln Leu Thr Met Glu Ser 340 345 350
- Asn Gly Lys Ser Val His Arg Asp Gly Ser Pro Val Ser Thr Gly Thr 355 360 365
- Gly Glu Ile Tyr Trp Gly Glu Pro Gly Thr Asn Gly Gln His Ala Phe 370 380
- Phe Gln Leu Ile His Gln Gly Thr Arg Leu Val Pro Ala Asp Phe Ile 385 390 395 400
- Gly Phe Ala Arg Pro Lys Gln Asp Leu Pro Ala Gly Glu Arg Thr Met 405 410 415
- His Asp Leu Leu Met Ser Asn Phe Phe Ala Gln Thr Lys Val Leu Ala 420 425 430
- Phe Gly Lys Asn Ala Glu Glu Ile Ala Ala Glu Gly Val Ala Pro Glu 435 440 445
- Leu Val Asn His Lys Val Met Pro Gly Asn Arg Pro Thr Thr Thr Ile 450 455 460
- Leu Ala Glu Glu Leu Thr Pro Ser Ile Leu Gly Ala Leu Ile Ala Leu 465 470 475 480

Tyr Glu His Ile Val Met Val Gln Gly Val Ile Trp Asp Ile Asn Ser 485 Phe Asp Gln Trp Gly Val Glu Leu Gly Lys Gln Gln Ala Asn Asp Leu 505 Ala Pro Ala Val Ser Gly Glu Glu Asp Val Asp Ser Gly Asp Ser Ser 520 Thr Asp Ser Leu Ile Lys Trp Tyr Arg Ala Asn Arg 535 <210> 43 <211> 630 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (1)..(630) <223> RXA01989 <400> 43 gtt aaa tca att cac aaa aca att cat gaa ggt act ggt gca ggt agt 48 Val Lys Ser Ile His Lys Thr Ile His Glu Gly Thr Gly Ala Gly Ser 1 15 gac ttc tta ggc tgg gtt gat tta cca gtt gat tac gac aaa gaa gaa Asp Phe Leu Gly Trp Val Asp Leu Pro Val Asp Tyr Asp Lys Glu Glu 20 30 144 ttt tca aga att gtt gaa gca tca aaa cgc att aaa gaa aat tct gat Phe Ser Arg Ile Val Glu Ala Ser Lys Arg Ile Lys Glu Asn Ser Asp 35 192 gtt tta gta gtc atc ggt att ggt ggt tct tac tta ggt gca cgt gca Val Leu Val Val Ile Gly Ile Gly Gly Ser Tyr Leu Gly Ala Arg Ala gca atc gaa atg tta acg tca tca ttt aga aac agc aat gaa tac cct Ala Ile Glu Met Leu Thr Ser Ser Phe Arg Asn Ser Asn Glu Tyr Pro 70 75 288 gaa att gta ttt gtt ggt aat cac tta tca tca aca tat acg aaa gag Glu Ile Val Phe Val Gly Asn His Leu Ser Ser Thr Tyr Thr Lys Glu 90 tta gtt gat tat tta gca gac aaa gat ttc tct gta aac gtt att tct 336 Leu Val Asp Tyr Leu Ala Asp Lys Asp Phe Ser Val Asn Val Ile Ser 105 aaa tot ggt aca act aca gaa cca gca gtt gca ttt aga ttg ttc aaa 384 Lys Ser Gly Thr Thr Glu Pro Ala Val Ala Phe Arg Leu Phe Lys 115 120 caa tta gtt gaa gaa aga tac ggt aaa gaa gaa gca caa aaa cgt ata 432 Gin Leu Val Glu Glu Arg Tyr Gly Lys Glu Glu Ala Gln Lys Arg Ile 130 135 140 ttt gca aca acg gat aaa gaa aaa ggt gct tta aaa cag ttg gct aca 480

Phe Ala Thr Thr Asp Lys Glu Lys Gly Ala Leu Lys Gln Leu Ala Thr 150 aac gaa ggt tat gaa acg ttt atc gta cct gat gat gta ggt gga aga 528 Asn Glu Gly Tyr Glu Thr Phe Ile Val Pro Asp Asp Val Gly Gly Arg 170 tat tot gtt tta aca gca gta gga tta tta cca att gca aca gct gga 576 Tyr Ser Val Leu Thr Ala Val Gly Leu Leu Pro Ile Ala Thr Ala Gly att aac atc gaa gct atg atg att ggt gct gca aaa gca cgt gaa gaa 624 Ile Asn Ile Glu Ala Met Met Ile Gly Ala Ala Lys Ala Arg Glu Glu 200 tta tct 630 Leu Ser 210 <210> 44 <211> 210 <212> PRT <213> Corynebacterium glutamicum <400> 44 Val Lys Ser Ile His Lys Thr Ile His Glu Gly Thr Gly Ala Gly Ser 10 Asp Phe Leu Gly Trp Val Asp Leu Pro Val Asp Tyr Asp Lys Glu Glu Phe Ser Arg Ile Val Glu Ala Ser Lys Arg Ile Lys Glu Asn Ser Asp Val Leu Val Val Ile Gly Ile Gly Gly Ser Tyr Leu Gly Ala Arg Ala Ala Ile Glu Met Leu Thr Ser Ser Phe Arg Asn Ser Asn Glu Tyr Pro Glu Ile Val Phe Val Gly Asn His Leu Ser Ser Thr Tyr Thr Lys Glu Leu Val Asp Tyr Leu Ala Asp Lys Asp Phe Ser Val Asn Val Ile Ser 100 Lys Ser Gly Thr Thr Thr Glu Pro Ala Val Ala Phe Arg Leu Phe Lys 120 Gln Leu Val Glu Glu Arg Tyr Gly Lys Glu Glu Ala Gln Lys Arg Ile 130 Phe Ala Thr Thr Asp Lys Glu Lys Gly Ala Leu Lys Gln Leu Ala Thr 155 Asn Glu Gly Tyr Glu Thr Phe Ile Val Pro Asp Asp Val Gly Gly Arg 170 Tyr Ser Val Leu Thr Ala Val Gly Leu Leu Pro Ile Ala Thr Ala Gly 180 185

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acc Thr	gaa Glu	acc Thr	aaa Lys	ccc Pro 170	acc Thr	aac Asn	tgg Trp	aac Asn	ggc Gly 175	gca Ala	acc Thr	aca Thr	gat Asp	ccc Pro 180	act Thr	643
cgt Arg	ttc Phe	ttg Leu	ttg Leu 185	ctt Leu	cgć Arg	cac His	ggc	caa Gln 190	act Thr	gct Ala	atg Met	tca Ser	gtg Val 195	gca Ala	cgc Arg	691
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caa Gln	gca Ala 215	gca Ala	gcg Ala	gca Ala	gca Ala	cga Arg 220	cga Arg	ctc Leu	gct Ala	caa Gln	acc Thr 225	ggt Gly	ggc Gly	atc Ile	gac Asp	787
gct Ala 230	att Ile	gtg Val	agt Ser	tct Ser	ccg Pro 235	ctc Leu	acc Thr	cgc Arg	acg Thr	atg Met 240	caa Gln	acc Thr	gca Ala	gaa Glu	gca Ala 245	835
gca Ala	gcg Ala	gcc Ala	gca Ala	ctg Leu 250	gga Gly	atg Met	aaa Lys	gta Val	cgt Arg 255	gtt Val	atc Ile	gat Asp	gat Asp	ctc Leu 260	atc Ile	883
gaa Glu	act Thr	gac Asp	ttt Phe 265	gga Gly	ctg Leu	tgg Trp	gat Asp	gga Gly 270	aaa Lys	tca Ser	ttt Phe	tca Ser	gaa Glu 275	gcc Ala	cac His	931
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ttc Phe	aac Asn 375	gac Asp	acc Thr	tca Ser	cac His	ctg Leu 380	gaa Glu	gcg Ala	tgac	gaca	gt c	tgac	ggaa	g		1266
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Leu Lys Glu Ile Ala Tyr Val Val Gly Thr Lys Ala Thr Asn Asn Val 35 40 45

Ala Glu Tyr Arg Gly Leu Leu Glu Gly Leu Lys Ala Ala Arg Glu Leu 50 55 60

Gly Ala Thr Ser Val Asp Val Tyr Met Asp Ser Lys Leu Val Val Glu 65 70 75 80

Gln Met Ser Gly Arg Trp Lys Ile Lys His Pro Asp Met Lys Val Leu 85 90 95

Ala Ile Glu Ala Lys Glu Ile Ala Ser Glu Ile Gly Ser Val Ser Tyr 100 105 110

Thr Trp Ile Pro Arg Glu Lys Asn Lys Arg Ala Asp Ala Leu Ser Asn 115 120 125

Val Ala Met Asp Ala Ala Ala Gly Lys Pro Val Gly Val Val Gly 130 135 140

Asp Ser Ala Ser Val Ser Ser Ala Ser Ser Val Ala Gly Ser Glu Lys 145 150 155 160

Glu Asp Leu Asn Cys Thr Glu Thr Lys Pro Thr Asn Trp Asn Gly Ala 165 170 175

Thr Thr Asp Pro Thr Arg Phe Leu Leu Leu Arg His Gly Gln Thr Ala 180 185 190

Met Ser Val Ala Arg Leu Tyr Ser Gly Arg Ser Asn Pro Glu Leu Ser 195 200 205

Glu Leu Gly Glu Lys Gln Ala Ala Ala Ala Ala Arg Arg Leu Ala Gln 210 215 220

Thr Gly Gly Ile Asp Ala Ile Val Ser Ser Pro Leu Thr Arg Thr Met 225 230 235 240

Gln Thr Ala Glu Ala Ala Ala Ala Ala Leu Gly Met Lys Val Arg Val 245 250 . 255

Ile Asp Asp Leu Ile Glu Thr Asp Phe Gly Leu Trp Asp Gly Lys Ser 260 265 270

Phe Ser Glu Ala His Glu Gln Asp Pro Glu Leu His Thr Lys Trp Leu 275 280 285

Thr Asp Ser Ser Val Ala Pro Pro Gly Gly Glu Ser Leu Gln Thr Val 295 Asn Arg Arg Val Lys Lys Ala Arg Glu Ser Leu Gln Arg Glu Tyr Gly Ala Ala Asn Val Leu Val Val Ser His Val Thr Pro Ile Lys Ala Ile 325 330 335 Met Arg Gln Ala Leu Asp Ala Gly Pro Ser Phe Phe Gln Lys Ala His 345 Leu Asp Leu Ala Ser Leu Ser Ile Ala Glu Phe Tyr Glu Asp Gly Pro 355 360 Thr Cys Val Arg Leu Phe Asn Asp Thr Ser His Leu Glu Ala 375 <210> 47 <211> 840 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(817) <223> RXA02492 <400> 47 gctgtacaac gacgctattg ccaacgaaaa tgtcgacggt gaaacgcatc acggctaagt 60 aaacgcgcgt cgtggaacat aaagtggcaa actagtacct atg act aac gga aaa Met Thr Asn Gly Lys ttg att ctt ctt cgt cac ggt cag agc gaa tgg aac gca tcc aac cag 163 Leu Ile Leu Leu Arg His Gly Gln Ser Glu Trp Asn Ala Ser Asn Gln ttc act gga tgg gtc gac gtc aat ctg acc gaa cag ggt gag gct gag Phe Thr Gly Trp Val Asp Val Asn Leu Thr Glu Gln Gly Glu Ala Glu gcc aaa ggc gtc ctc cca ggc gtt gta tac acc tcc ttg ctg cgc 259 Ala Lys Gly Val Leu Pro Gly Val Val Tyr Thr Ser Leu Leu Arg Arg 40 gcg atc cgc act gca aac atc gca ctg aac gct gca gac cgc cac tgg 307 Ala Ile Arg Thr Ala Asn Ile Ala Leu Asn Ala Ala Asp Arg His Trp 55 60 atc cca gtg atc cgc gac tgg cgc ctc aac gag cgt cac tac ggc gca 355 Ile Pro Val Ile Arg Asp Trp Arg Leu Asn Glu Arg His Tyr Gly Ala 70 ctg cag ggc ctt gac aag gct gca acc aag gaa aaa tac ggc gac gac Leu Gln Gly Leu Asp Lys Ala Ala Thr Lys Glu Lys Tyr Gly Asp Asp 95

cag ttc atg gaa tgg cgc cgc tcc tac gac acc cca cca cca gag ctc Gln Phe Met Glu Trp Arg Arg Ser Tyr Asp Thr Pro Pro Pro Glu Leu 105 110 115	451
gcg gat gac gca gag tac tcc cag gca aat gac cct cgt tac gcg gac Ala Asp Asp Ala Glu Tyr Ser Gln Ala Asn Asp Pro Arg Tyr Ala Asp 120 125 130	499
ctc gac gta gtt cca cgc acc gaa tgc ctc aag gac gtt gtg gtt cgt Leu Asp Val Val Pro Arg Thr Glu Cys Leu Lys Asp Val Val Val Arg 135 140 145	547
ttt gtt cct tac ttc gag gaa gaa atc ctg cca cgc gca aag aag ggc Phe Val Pro Tyr Phe Glu Glu Glu Ile Leu Pro Arg Ala Lys Lys Gly 150 155 160 165	7
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aag cac ctt gac ggc atc tcc gat gct gat atc gca gag ctc aac atc Lys His Leu Asp Gly Ile Ser Asp Ala Asp Ile Ala Glu Leu Asn Ile 185 190 195	
cca acc ggc atc cca ctg gtc tac gaa atc gcc gaa gac ggt tcc gta Pro Thr Gly Ile Pro Leu Val Tyr Glu Ile Ala Glu Asp Gly Ser Val 200 205 210	
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Pro	Pro	Pro 115	Glu	Leu	Ala	Asp	Asp 120		Glu	Tyr	Ser	Gln 125		Asn	Asp	
Pro	Arg 130	Tyr	Ala	Asp	Leu	Asp 135		Val	Pro	Arg	Thr 140	Glu	Cys	Leu	Lys	
Asp 145	Val	Val	Val	Arg	Phe 150		Pro	Tyr	Phe	Glu 155		Glu	Ile	Leu	Pro 160	
Arg	Ala	Lys	Lys	Gly 165	Glu	Thr	Val	Leu	Ile 170	Ala	Ala	His	Gly	Asn 175		
Leu	Arg	Ala	Leu 180	Val	Lys	His	Leu	Asp 185	Gly	Ile	Ser	Asp	Ala 190	Asp	Ile	
Ala	Glu	Leu 195	Asn	Ile	Pro	Thr	Gly 200	Ile	Pro	Leu	Val	Tyr 205	Glu	Ile	Ala	
Glu	Asp 210	Gly	Ser	Val	Val	Asn 215	Pro	Gly	Gly	Thr	Tyr 220	Leu	Asp	Pro	Glu	
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gtc Val	cat His	cta Leu	gtt Val	cgc Arg 10	cac His	ggc Gly	gaa Glu	gtc Val	cac His 15	aac Asn	cca Pro	gag Glu	aaa Lys	atc Ile 20	ctg Leu	163
tac Tyr	gga Gly	cgc Arg	atg Met 25	ccc Pro	gga Gly	tac Tyr	agg Arg	ttg Leu 30	tct Ser	tcc Ser	cgt Arg	gga Gly	cgc Arg 35	agc Ser	caa Gln	211
gcc Ala	gcc Ala	cgc Arg	act Thr	gca Ala	gct Ala	tct Ser	ttt Phe	gaa Glu	ggc Gly	cac His	gat Asp	gtc Val	acc Thr	tac Tyr	att Ile	259

307

gcg gcc tcc cca ttg cag cgt gtg cag gaa acc tcc gaa ccg ttc atc

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40

60 55 65 355 aag gtc aca ggc cta gaa ctg atc acc gac gag gat ctt ctg gaa gca Lys Val Thr Gly Leu Glu Leu Ile Thr Asp Glu Asp Leu Leu Glu Ala 75 80 ggc aac cgt ttc gaa ggc ctg cgc acc aaa ggt tgg cgt tcc cag ttg 403 Gly Asn Arg Phe Glu Gly Leu Arg Thr Lys Gly Trp Arg Ser Gln Leu 95 90 tgg aac ccc gtg cgt tgg cct ttg atg tac aac ccc acg ctt ccc agc 451 Trp Asn Pro Val Arg Trp Pro Leu Met Tyr Asn Pro Thr Leu Pro Ser 110 499 tgg ggc gaa cac tac acc gac att ttg gaa aga atg atg gcg gct gtg Trp Gly Glu His Tyr Thr Asp Ile Leu Glu Arg Met Met Ala Ala Val 120 125 130 gaa cga gct cgg gtg gca gcg gaa gga cac gaa gca atc ctg gtg acc 547 Glu Arg Ala Arg Val Ala Ala Glu Gly His Glu Ala Ile Leu Val Thr 135 140 595 cac cag ttg ccg atc gtg tgc gtg caa cgc cac gcc cgc gga caa agc His Gln Leu Pro Ile Val Cys Val Gln Arg His Ala Arg Gly Gln Ser 160 150 155 ctg tcc cat aac cca gcg acc agg caa tgc gac ctc gcc tca gtg aca 643 Leu Ser His Asn Pro Ala Thr Arg Gln Cys Asp Leu Ala Ser Val Thr 180 170 175 tcc ttg gtg ttc caa gac gat caa att gtc ggc gtg cat tac aac gaa Ser Leu Val Phe Gln Asp Asp Gln Ile Val Gly Val His Tyr Asn Glu 185 190 729 cca gct cag gag att tgatcactcg tgcgtttgac caa Pro Ala Gln Glu Ile 200 <210> 50 <211> 202 <212> PRT <213> Corynebacterium glutamicum <400> 50 Met Thr Gln Thr Ile Val His Leu Val Arg His Gly Glu Val His Asn Pro Glu Lys Ile Leu Tyr Gly Arg Met Pro Gly Tyr Arg Leu Ser Ser Arg Gly Arg Ser Gln Ala Ala Arg Thr Ala Ala Ser Phe Glu Gly His Asp Val Thr Tyr Ile Ala Ala Ser Pro Leu Gln Arg Val Gln Glu Thr 50

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Asp Leu Leu Glu Ala Gly Asn Arq Phe Glu Gly Leu Arg Thr Lys Gly

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tta Leu	ctc Leu	att Ile	cgg Arg 25	cat His	Gly	caa Gln	acc Thr	cca Pro 30	aca Thr	act Thr	ggt Gly	cag Gln	gtt Val 35	ctg Leu	cct Pro	211
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cgg Arg	gag Glu 55	gtg Val	gca Ala•	cag Gln	cgt Arg	ctg Leu 60	gcg Ala	gag Glu	gtg Val	ccg Pro	att Ile 65	acc Thr	gct Ala	gtg Val	tat Tyr	307
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gct cat ggc Ala His Gly												403	
ttc ggc gag Phe Gly Glu												451	
gag tgg aaa Glu Trp Lys 120	Ala Val			-								499	
ggt gag agt Gly Glu Ser 135			Gln									547	
aac att gcg Asn Ile Ala 150												595	
gcc gac acg Ala Asp Thr												643	
gat tct ttt Asp Ser Phe			Ile	-	_					_		691	
gaa ttt acc Glu Phe Thr 200												739	
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Gly Gln Val	Leu Pro	Gly Gln	Thr 1	Pro	Gly	Leu	His	Leu 45	Ser	Asp	Lys		
Gly Glu Glu 50	Gln Ala	Arg Glu 55	Val a	Ala	Gln	Arg	Leu 60	Ala	Glu	Val	Pro		
Ile Thr Ala 65	Val Tyr	Ser Ser 70	Pro i	Met	Glu	Arg 75	Ala	Gln	Glu	Thr	Ala 80		

Ala	Pro	Thr	Val	Ser 85		His	Gly	Leu	Glu 90	Leu	Thr	Val	Glu	Pro 95	Gly	
Leu	Ile	Glu	Cys 100	Asp	Phe	Gly	Glu	Trp 105		Gly	Arg	Lys	Leu 110		Glu	
Leu	Asn	Ala 115	Leu	Glu	Glu	Trp	Lys 120		Val	Gln	Lys	Thr 125		Ser	Thr	
Phe	Arg 130	Phe	Pro	Gly	Gly	Glu 135		Phe	Val	Glu	Met 140	Gln	Asp	Arg	Met	
Val 145	Glu	Ala	Ile	Gly	Asn 150	Ile	Ala	Gln	Gln	His 155	Pro	Gly	Glu	Ile	Val 160	
Ala	Ala	Phe	Ser	His 165	Ala	Asp	Thr	Ile	Lys 170	Ala	Ala	Val	Ala	His 175	Phe	
Val	Gly	Thr	Pro 180	Leu	Asp	Ser	Phe	Gln 185	Arg	Ile	Phe	Ile	Asp 190	Thr	Ala	
Ser	Ile	Ser 195	Ala	Val	Glu	Phe	Thr 200	Gly	Lys	Ser	Ser	Gly 205	Val	Ser	Ser	
His	Met 210	Leu	Leu	Thr	Asn	Ser 215	Arg	Thr	Gly	Ser	Leu 220	Gly	Tyr	Leu	Arg	
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	gtgat									aat		gaa	gac	atg	cga	115
att Ile	gct Ala	act Thr	ctc Leu	acg Thr 10	tca Ser	ggc Gly	ggc Gly	gac Asp	tgc Cys 15	ccc Pro	gga Gly	cta Leu	aac Asn	gcc Ala 20	gtc Val	163
atc Ile	cga Arg	gga Gly	atc Ile 25	gtc Val	cgc Arg	aca Thr	gcc Ala	agc Ser 30	aat Asn	gaa Glu	ttt Phe	ggc Gly	tcc Ser 35	acc Thr	gtc Val	211
gtt Val	ggt Gly	tat Tyr (caa Gln	gac Asp	ggt Gly	tgg Trp	gaa Glu 45	gga Gly	ctg Leu	tta Leu	ggc Gly	gat Asp 50	cgt Arg	cgc Arg	gta Val	259

						gat Asp 60										307
						cgc Arg										355
						aac Asn										403
					_	gga Gly		_	_		_	_		_		451
						gtc Val										499
						acc Thr 140										547
						cgc Arg										595
						gtc Val										643
						ggc										691
						gag Glu										739
						ggc Gly 220										787 _.
	_	-			_	gag Glu		_	-				-	_		835
						gga Gly										883
						gat Asp										931
						act Thr										979
tat	ggt	gtt	cgt	gca	gct	cgt	gcg	tgc	cat	gag	gga	agc	ttt	gac	aag	1027

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<213> Corynebacterium glutamicum

<400> 54

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Gly Leu Asn Ala Val Ile Arg Gly Ile Val Arg Thr Ala Ser Asn Glu 20 25 30

Phe Gly Ser Thr Val Val Gly Tyr Gln Asp Gly Trp Glu Gly Leu Leu 35 40 45

Gly Asp Arg Arg Val Gln Leu Tyr Asp Asp Glu Asp Ile Asp Arg Ile
50 55 60

Leu Leu Arg Gly Gly Thr Ile Leu Gly Thr Gly Arg Leu His Pro Asp 65 70 75 80

Lys Phe Lys Ala Gly Ile Asp Gln Ile Lys Ala Asn Leu Glu Asp Ala 85 90 95

Gly Ile Asp Ala Leu Ile Pro Ile Gly Gly Glu Gly Thr Leu Lys Gly
100 105 110

Ala Lys Trp Leu Ser Asp Asn Gly Ile Pro Val Val Gly Val Pro Lys 115 120 125

Thr Ile Asp Asn Asp Val Asn Gly Thr Asp Phe Thr Phe Gly Phe Asp 130 135 140

Thr Ala Val Ala Val Ala Thr Asp Ala Val Asp Arg Leu His Thr Thr 145 150 155 160

Ala Glu Ser His Asn Arg Val Met Ile Val Glu Val Met Gly Arg His
165 170 175

Val Gly Trp Ile Ala Leu His Ala Gly Met Ala Gly Gly Ala His Tyr 180 185 190

Thr Val Ile Pro Glu Val Pro Phe Asp Ile Ala Glu Ile Cys Lys Ala 195 200 205

Met Glu Arg Arg Phe Gln Met Gly Glu Lys Tyr Gly Ile Ile Val Val 215 Ala Glu Gly Ala Leu Pro Arg Glu Gly Thr Met Glu Leu Arg Glu Gly 235 His Ile Asp Gln Phe Gly His Lys Thr Phe Thr Gly Ile Gly Gln Gln 250 Ile Ala Asp Glu Ile His Val Arg Leu Gly His Asp Val Arg Thr Thr Val Leu Gly His Ile Gln Arg Gly Gly Thr Pro Thr Ala Phe Asp Arg 280 Val Leu Ala Thr Arg Tyr Gly Val Arg Ala Ala Arg Ala Cys His Glu Gly Ser Phe Asp Lys Val Val Ala Leu Lys Gly Glu Ser Ile Glu Met 315 Ile Thr Phe Glu Glu Ala Val Gly Thr Leu Lys Glu Val Pro Phe Glu 325 Arg Trp Val Thr Ala Gln Ala Met Phe Gly 340 <210> 55 <211> 1083 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1060) <223> RXA01243 <400> 55 gcgcaatcag cgatatcgat gtggtggtca ccgatgcggg tgcaccagca agtttcgttg 60 agcagttgcg agaacgcgat gtagaagttg tgattgcaga atg att ctt aca gtc Met Ile Leu Thr Val 1 act gca agt ccg tat ctg ttg agc acc aat gag ctt gac ggc acc atc Thr Ala Ser Pro Tyr Leu Leu Ser Thr Asn Glu Leu Asp Gly Thr Ile 10 211 gaa att ggc gaa gca aac aaa atc cgg cag gtt tcc act gtt gcc ggt Glu Ile Gly Glu Ala Asn Lys Ile Arg Gln Val Ser Thr Val Ala Gly 25 ggt ttt ggc acc ggt gtg gct gcc acc ttg ttt tat ggc ggc aat gaa Gly Phe Gly Thr Gly Val Ala Ala Thr Leu Phe Tyr Gly Gly Asn Glu 40 45 act ttt gca gtt ttt ccc gct cca gaa atc tct cat tac atg cgc ctg Thr Phe Ala Val Phe Pro Ala Pro Glu Ile Ser His Tyr Met Arg Leu 55

														ggt Gly		355
														act Thr 100		403
														att Ile		451
														ttg Leu		499
ggt Gly	ggc Gly 135	aat Asn	ttg Leu	ccg Pro	tct Ser	atc Ile 140	gcg Ala	cct Pro	gct Ala	gcg Ala	tgg Trp 145	ttt Phe	gtg Val	gat Asp	gtg Val	547
														atc Ile		595
gca Ala	act Thr	ggt Gly	gct Ala	gcg Ala 170	ttg Leu	cgt Arg	gcg Ala	gtt Val	att Ile 175	cga Arg	cag Gln	ctt Leu	gca Ala	gct Ala 180	acg Thr	643
tcc Ser	ccg Pro	gat Asp	gcg Ala 185	ctg Leu	att Ile	gtg Val	gct Ala	gcg Ala 190	gaa Glu	gaa Glu	atc Ile	gaa Glu	att Ile 195	gcc Ala	act Thr	691
														gat Asp		739
														gtc Val		787
gag Glu 230	gtg Val	ttg Leu	gtt Val	acc Thr	aac Asn 235	aag Lys	cgg Arg	acg Thr	gaa Glu	tct Ser 240	ttg Leu	tat Tyr	gtt Val	tcc Ser	gag Glu 245	835
tct Ser	gaa Glu	tca Ser	ctg Leu	tta Leu 250	gcc Ala	agc Ser	tac Tyr	gac Asp	agc Ser 255	acc Thr	cct Pro	ggt Gly	aag Lys	cag Gln 260	ggc Gly	883
gtg Val	aat Asn	tgg Trp	cgg Arg 265	gaa Glu	tct Ser	ttt Phe	act Thr	gca Ala 270	gga Gly	ttc Phe	ttg Leu	gca Ala	gca Ala 275	tcc Ser	aat Asn	931
														tac Tyr		979
aac Asn	gct Ala 295	gaa Glu	ggc Gly	agt Ser	gag Glu	tgg Trp 300	gac Asp	aac Asn	tac Tyr	att Ile	ccc Pro 305	aca Thr	ccc Pro	gat Asp	aag Lys	1027
ctt	cgg	gcg	gag	cac	gtg	gtc	atc	aaa	tcg	ctt	taga	ccac	gc a	aaaa	gcctc	1080

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aaa 1083

<210> 56

<211> 320

<212> PRT

<213> Corynebacterium glutamicum

<400> 56

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Leu Asp Gly Thr Ile Glu Ile Gly Glu Ala Asn Lys Ile Arg Gln Val 20 25 30

Ser Thr Val Ala Gly Gly Phe Gly Thr Gly Val Ala Ala Thr Leu Phe 35 40 45

Tyr Gly Gly Asn Glu Thr Phe Ala Val Phe Pro Ala Pro Glu Ile Ser 50 60

His Tyr Met Arg Leu Val Thr Phe Ala Gly Leu Pro His Glu Ile Ile 65 70 75 80

Pro Val Ala Gly Pro Ile Pro Met His Leu Thr Met Arg Asp Ala Glu 85 90 95

Gly Asn Glu Thr Lys Phe Lys Asp Ser Pro Met Pro Leu Asp Val Ser 100 105 110

Gln Leu Ala Ile Leu Arg Asp Leu Val Val Arg Arg Ala Glu Asp Ala 115 120 125

Ala Trp Val Leu Leu Gly Gly Asn Leu Pro Ser Ile Ala Pro Ala Ala 130 135 140

Trp Phe Val Asp Val Val Arg Ser Leu Arg Leu Tyr His Pro His Val
145 150 155 160

Lys Val Ala Ile Ala Ala Thr Gly Ala Ala Leu Arg Ala Val Ile Arg 165 170 175

Gln Leu Ala Ala Thr Ser Pro Asp Ala Leu Ile Val Ala Ala Glu Glu 180 185 190

Ile Glu Ile Ala Thr Gly Leu Glu Pro Lys Thr Leu Arg Gly Pro Trp 195 200 205

Val Glu Gly Asp Leu Ser Pro Thr Val Ala Ala Ala Arg Ala Leu Ile 210 215 220

Asp Ser Gly Val Thr Glu Val Leu Val Thr Asn Lys Arg Thr Glu Ser 225 230 235 240

Leu Tyr Val Ser Glu Ser Glu Ser Leu Leu Ala Ser Tyr Asp Ser Thr 245 250 255

Pro Gly Lys Gln Gly Val Asn Trp Arg Glu Ser Phe Thr Ala Gly Phe

260 265 270

Leu Ala Ala Ser Asn Asp Gly Lys Ser Thr Glu Asp Ser Val Ile Asn 275 280 285

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Pro Thr Pro Asp Lys Leu Arg Ala Glu His Val Val Ile Lys Ser Leu 305 310 315 320

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451

ctc aac ggc ccc ggc gcg ccg ctc agc gag cag aag ctc cgt agc ttg

Leu Asn Gly Pro Gly Ala Pro Leu Ser Glu Gln Lys Leu Arg Ser Leu

gaa aag gtg ctt atc gac gcg ctc cgc ccc gaa gtc acc tgg gtt gtc

110

105

PCT/IB00/00943 WO 01/00844

Glu Lys	Val Leu 120	Ile Asp	Ala	Leu 125	Arg	Pro	Glu	Val	Thr 130	Trp	Val	Val	
ctg gcg Leu Ala 135	ggc tcg Gly Ser	ctg cca Leu Pro	cca Pro 140	Gly ggg	gca Ala	cca Pro	gtt Val	gac Asp 145	tgg Trp	tac Tyr	gcg Ala	cgt Arg	547
	gcg ttg Ala Leu		Ser										595
gat acc Asp Thr	tca gac Ser Asp	aag cca Lys Pro	ctg Leu	atg Met	gcg Ala	ttg Leu 175	ggc Gly	gag Glu	agc Ser	ttg Leu	gat Asp 180	aca Thr	643
cct ggc Pro Gly	gct gct Ala Ala 185	ccg aad Pro Asr	ctg Leu	att Ile	aag Lys 190	cca Pro	aat Asn	ggt Gly	ctg Leu	gaa Glu 195	ctg Leu	ggc Gly	691
cag ctg Gln Leu	gct aac Ala Asn 200	act gat Thr Asp	ggt Gly	gaa Glu 205	gag Glu	ctg Leu	gag Glu	gcg Ala	cgt Arg 210	gct Ala	gcg Ala	caa Gln	739
ggc gat Gly Asp 215	tac gac Tyr Asp	gcc ato Ala Ile	atc Ile 220	gca Ala	gct Ala	gcg Ala	gac Asp	gta Val 225	ctg Leu	gtt Val	aac Asn	cgt Arg	787
ggc atc Gly Ile 230	gaa cag Glu Gln	gtg ctt Val Leu 235	. Val	acc Thr	ttg Leu	ggt Gly	gcc Ala 240	gca Ala	gga Gly	gcg Ala	gtg Val	ttg Leu 245	835
gtc aac Val Asn	gca gaa Ala Glu	ggt gcg Gly Ala 250	tgg Trp	act Thr	gct Ala	act Thr 255	tct Ser	cca Pro	aag Lys	att Ile	gat Asp 260	gtt Val	883
gta tcc Val Ser	acc gtt Thr Val 265	gga gct Gly Ala	gga Gly	gac Asp	tgt Cys 270	gct Ala	ctt Leu	gca Ala	ggt Gly	ttt Phe 275	gtt Val	atg Met	931
gca cgt Ala Arg	tcc cag Ser Gln 280	aag aaa Lys Lys	aca Thr	ctg Leu 285	gag Glu	gaa Glu	tct Ser	ctg Leu	ctg Leu 290	aat Asn	gcc Ala	gtg Val	979
tct tac Ser Tyr 295	ggc tcg Gly Ser	act gcc Thr Ala	gcg Ala 300	tct Ser	ctt Leu	cct Pro	ggc Gly	act Thr 305	acc Thr	att Ile	cct Pro	cgt Arg	1027
cct gac Pro Asp 310	caa ctc Gln Leu	gcc aca Ala Thr 315	Ala	ggt Gly	gca Ala	acg Thr	gtc Val 320	acc Thr	caa Gln	gtc Val	aaa Lys	gga Gly 325	1075
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<400> 58

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Leu Ala Gly Phe Glu Thr Leu Ala Val Phe Pro Ala Gly Lys Leu Asp 50 55 60

Pro Phe Val Pro Leu Val Arg Asp Ile Gly Leu Pro Val Glu Thr Val 65 70 75 80

Val Ile Asn Lys Asn Val Arg Thr Asn Thr Thr Val Thr Glu Pro Asp 85 90 95

Gly Thr Thr Lys Leu Asn Gly Pro Gly Ala Pro Leu Ser Glu Gln
100 105 110

Lys Leu Arg Ser Leu Glu Lys Val Leu Ile Asp Ala Leu Arg Pro Glu
115 120 125

Val Thr Trp Val Val Leu Ala Gly Ser Leu Pro Pro Gly Ala Pro Val 130 135 140

Asp Trp Tyr Ala Arg Leu Thr Ala Leu Ile His Ser Ala Arg Pro Asp 145 150 155 160

Val Arg Val Ala Val Asp Thr Ser Asp Lys Pro Leu Met Ala Leu Gly
165 170 175

Glu Ser Leu Asp Thr Pro Gly Ala Ala Pro Asn Leu Ile Lys Pro Asn 180 185 190

Gly Leu Glu Leu Gly Gln Leu Ala Asn Thr Asp Gly Glu Glu Leu Glu 195 200 205

Ala Arg Ala Ala Gln Gly Asp Tyr Asp Ala Ile Ile Ala Ala Asp 210 215 220

Val Leu Val Asn Arg Gly Ile Glu Gln Val Leu Val Thr Leu Gly Ala 225 230 235 240

Ala Gly Ala Val Leu Val Asn Ala Glu Gly Ala Trp Thr Ala Thr Ser 245 250 255

Pro Lys Ile Asp Val Val Ser Thr Val Gly Ala Gly Asp Cys Ala Leu 260 265 270

Ála Gly Phe Val Met Ala Arg Ser Gln Lys Lys Thr Leu Glu Glu Ser 275 280 285

Leu Leu Asn Ala Val Ser Tyr Gly Ser Thr Ala Ala Ser Leu Pro Gly 290 295 300

Thr Thr Ile Pro Arg Pro Asp Gln Leu Ala Thr Ala Gly Ala Thr Val 305 310 315 320 Thr Gln Val Lys Gly Leu Lys Glu Ser Ala 325 330

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PCT/IB00/00943 WO 01/00844

ggt ggc gaa gaa Gly Gly Glu Glu	gac ggc gtt Asp Gly Val 170	gag gct aag Glu Ala Lys 175	gct ggc gca a Ala Gly Ala A	aac ctc tac Asn Leu Tyr 180	643
acc tcc cca gaa Thr Ser Pro Glu 185	gac ttt gag Asp Phe Glu	aag acc atc Lys Thr Ile 190	Asp Ala Ile (ggc acc ggt Gly Thr Gly 195	691
gag aag ggc cgc Glu Lys Gly Arg 200					739
gtt tac aag cca Val Tyr Lys Pro 215		. Lys Leu Arg			787
ggc cag cag gtt Gly Gln Gln Val 230					835
cca ttc gac ttc Pro Phe Asp Phe					883
atc gaa gag gcg Ile Glu Glu Ala 265			Lys Met Asn		931
gac acc cag tac Asp Thr Gln Tyr 280	gca ttc acc Ala Phe Th	c cgc cca atc Arg Pro Ile 285	gtc tcc cac Val Ser His 1 290	atg ttt gag Met Phe Glu	979
aac tac aac ggc Asn Tyr Asn Gly 295		Ile Asp Gly			1027
gct tac gac cca Ala Tyr Asp Pro 310					1075
gag cgc att atc Glu Arg Ile Ile	gag tct tge Glu Ser Cy: 330	c cag gac ctc s Gln Asp Leu 335	Lys Ser Val	gga aag acc Gly Lys Thr 340	1123
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- Ile Ile Gln Phe Ser Thr Gly Gly Ala Glu Phe Gly Ser Gly Leu Ala 50 55 60
- Val Lys Asn Lys Val Lys Gly Ala Val Ala Leu Ala Ala Phe Ala His 65 70 75 80
- Glu Ala Ala Lys Ser Tyr Gly Ile Asn Val Ala Leu His Thr Asp His 85 90 95
- Cys Gln Lys Glu Val Leu Asp Glu Tyr Val Arg Pro Leu Leu Ala Ile 100 105 110
- Ser Gln Glu Arg Val Asp Arg Gly Glu Leu Pro Leu Phe Gln Ser His 115 120 125
- Met Trp Asp Gly Ser Ala Val Pro Ile Asp Glu Asn Leu Glu Ile Ala 130 135 140
- Gln Glu Leu Leu Ala Lys Ala Lys Ala Ala Asn Ile Ile Leu Glu Val 145 150 155 160
- Glu Ile Gly Val Val Gly Gly Glu Glu Asp Gly Val Glu Ala Lys Ala 165 170 175
- Gly Ala Asn Leu Tyr Thr Ser Pro Glu Asp Phe Glu Lys Thr Ile Asp 180 185 190
- Ala Ile Gly Thr Gly Glu Lys Gly Arg Tyr Leu Leu Ala Ala Thr Phe 195 200 205
- Gly Asn Val His Gly Val Tyr Lys Pro Gly Asn Val Lys Leu Arg Pro 210 215 220
- Glu Val Leu Leu Glu Gly Gln Gln Val Ala Arg Lys Lys Leu Gly Leu 225 230 235 240
- Ala Asp Asp Ala Leu Pro Phe Asp Phe Val Phe His Gly Gly Ser Gly 245 250 255
- Ser Glu Lys Glu Lys Ile Glu Glu Ala Leu Thr Tyr Gly Val Ile Lys 260 265 270
- Met Asn Val Asp Thr Asp Thr Gln Tyr Ala Phe Thr Arg Pro Ile Val 275 280 285
- Ser His Met Phe Glu Asn Tyr Asn Gly Val Leu Lys Ile Asp Gly Glu 290 295 300
- Val Gly Asn Lys Lys Ala Tyr Asp Pro Arg Ser Tyr Met Lys Lys Ala 305 310 315 320
- Glu Gln Ser Met Ser Glu Arg Ile Ile Glu Ser Cys Gln Asp Leu Lys 325 330 335
- Ser Val Gly Lys Thr Thr Ser Lys 340

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ggc	cacc	ggg (aaac	tttt	tt a	agaa	aggt	g tg	tttc	acac					cca Pro 5	115
ctt Leu	atc Ile	gct Ala	ggt Gly	aac Asn 10	tgg Trp	aag Lys	atg Met	aac Asn	ctg Leu 15	gat Asp	cac His	cag Gln	cag Gln	gca Ala 20	atc Ile	163
							ttc Phe									211
							gtt Val 45									259
							aag Lys									307
gac Asp 70	gtc Val	tcc Ser	cag Gln	cac His	gag Glu 75	tcc Ser	ggt Gly	gcg Ala	tac Tyr	acc Thr 80	ggt Gly	gaa Glu	gtt Val	tct Ser	gca Ala 85	355
							tgc Cys									403
gag Glu	cgc Arg	cgc Arg	gag Glu 105	tac Tyr	cac His	aac Asn	gag Glu	tct Ser 110	gat Asp	gag Glu	ttg Leu	gtt Val	gct Ala 115	gcg Ala	aag Lys	451
							ggc Gly 125									499
							gct Ala									547
gag Glu 150	cag Gln	acc Thr	cgt Arg	aag Lys	tcc Ser 155	ctt Leu	gct Ala	ggc Gly	ctg Leu	gat Asp 160	gct Ala	gct Ala	gag Glu	ctg Leu	gcc Ala 165	595
aac Asn	acc Thr	gtt Val	atc Ile	gcg Ala 170	tat Tyr	gag Glu	cca Pro	gtg Val	tgg Trp 175	gct Ala	atc Ile	ggc Gly	acc Thr	ggt Gly 180	aag Lys	643

gtt gct Val Ala		Ala												691
ctg atc Leu Ile														739
ctt tac Leu Tyr 215														787
cag cct Gln Pro 230														835
gaa gca Glu Ala														877
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His Gln	Gln Ala 20		Gly	Thr	Val	Gln 25	Lys	Leu	Ala	Phe	Ala 30	Leu	Pro	
Lys Glu	Tyr Phe	Glu	Lys	Val	Asp 40	Val	Ala	Val	Thr	Val 45	Pro	Phe	Thr	
Asp Ile 50	Arg Ser	Val	Gln	Thr 55	Leu	Val	Glu	Gly	Asp 60	Lys	Leu	Glu	Val	
Thr Phe 65	Gly Ala	Gln	Asp 70	Val	Ser	Gln	His	Glu 75	Ser	Gly	Ala	Tyr	Thr 80	
Gly Glu	Val Ser	Ala 85	Ser	Met	Leu	Ala	Lys 90	Leu	Asn	Cys	Ser	Trp 95	Val	
Val Val	Gly His 100		Glu	Arg	Arg	Glu 105	Tyr	His	Asn	Glu	Ser 110	Asp	Glu	
Leu Val	Ala Ala 115	Lys	Ala	Lys	Ala 120	Ala	Leu	Ser	Asn	Gly 125	Ile	Ser	Pro	
Ile Val	Cys Val	Gly		Pro 135	Leu	Glu	Ile	Arg	Glu 140	Ala	Gly	Thr	His	
Val Glu 145	Tyr Val		Glu 150	Gln	Thr	Arg	Lys	Ser 155	Leu	Ala	Gly	Leu	Asp 160	
Ala Ala	Glu Leu	Ala 165	Asn	Thr	Val	Ile	Ala 170	Tyr	Glu	Pro	Val	Trp 175	Ala	

Ile Gly Thr Gly Lys Val Ala Ser Ala Ala Asp Ala Gln Glu Val Cys Lys Ala Ile Arg Gly Leu Ile Val Glu Leu Ala Gly Asp Glu Val Ala 200 Glu Gly Leu Arg Ile Leu Tyr Gly Gly Ser Val Lys Ala Glu Thr Val Ala Glu Ile Val Gly Gln Pro Asp Val Asp Gly Gly Leu Val Gly Gly 235 Ala Ser Leu Asp Gly Glu Ala Phe Ala Lys Leu Ala Ala Asn Ala Ala 245 250 Ser Val Ala <210> 63 <211> 1563 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1540) <223> RXN01225 <400> 63 tttgggctaa tgttgggggg agtgctttca actatccacg agagctgccc agtgataaac 60 cccgggttaa ccccacgcct aagtcagtga aggacttttt atg acg cac aac cac Met Thr His Asn His aag gac tgg aac gat cgc att gca gtt gcg gag gaa atg gtg ccg ttg Lys Asp Trp Asn Asp Arg Ile Ala Val Ala Glu Glu Met Val Pro Leu 10 atc ggg cgc ctg cac cgc aac aac atc gtg gtg gtt tcc gta ttc ggt Ile Gly Arg Leu His Arg Asn Asn Asn Val Val Val Ser Val Phe Gly 25 cgt ctc ctt gtg aat gtc tca gac atc gat atc atc aag tct cac cgc Arg Leu Leu Val Asn Val Ser Asp Ile Asp Ile Ile Lys Ser His Arg 40 50 tac gcc cgc cac atc ata tcc aag gaa ctt cca ctg gaa agc tcc ttg Tyr Ala Arg His Ile Ile Ser Lys Glu Leu Pro Leu Glu Ser Ser Leu 55 gat att ttg cgc gaa ctg gta gat atg aac ctt ggt acc gca tcg atc 355 Asp Ile Leu Arg Glu Leu Val Asp Met Asn Leu Gly Thr Ala Ser Ile 75 gac ctg gga cag ctg gcc tac agc ttc gaa gaa tcc gaa agc acc gac Asp Leu Gly Gln Leu Ala Tyr Ser Phe Glu Glu Ser Glu Ser Thr Asp 95

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						gcc Ala 155											595
						tcc Ser											643
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						cgc Arg 235											835
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	-				-	cgc Arg				-				_	_		1027
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acc acc cac Thr Thr His												1411
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Val Tyr Pro		Arg (_	-	Leu	tago	yttat	icc a	agco	ctaata	1560 1563
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Ser Glu Ser Thr Asp Leu Arg Ala Phe Leu Glu Asp Ala Leu Ala Pro 105 100 Val Ile Gly Ala Glu Thr Asp Ile Asn Pro Thr Asp Ile Val Leu Tyr 120 Gly Phe Gly Arg Ile Gly Arg Leu Leu Ala Arg Ile Leu Val Ser Arg Glu Ala Leu Tyr Asp Gly Ala Arg Leu Arg Ala Ile Val Val Arg Lys 150 155 Asn Gly Glu Glu Asp Leu Val Lys Arg Ala Ser Leu Leu Arg Arg Asp Ser Val His Gly Gly Phe Asp Gly Thr Ile Thr Thr Asp Tyr Asp Asn 185 Asn Ile Ile Trp Ala Asn Gly Thr Pro Ile Lys Val Ile Tyr Ser Asn Asp Pro Ala Thr Ile Asp Tyr Thr Glu Tyr Gly Ile Asn Asp Ala Val Val Val Asp Asn Thr Gly Arg Trp Arg Asp Arg Glu Gly Leu Ser Gln 235 His Leu Lys Ser Lys Gly Val Ala Lys Val Val Leu Thr Ala Pro Gly Lys Gly Asp Leu Lys Asn Ile Val Tyr Gly Ile Asn His Thr Asp Ile Thr Ala Asp Asp Gln Ile Val Ser Ala Ala Ser Cys Thr Thr Asn Ala 275 Ile Thr Pro Val Leu Lys Val Ile Asn Asp Arg Tyr Gly Val Glu Phe 295 Gly His Val Glu Thr Val His Ser Phe Thr Asn Asp Gln Asn Leu Ile 305 Asp Asn Phe His Lys Gly Ser Arg Arg Gly Arg Ala Ala Gly Leu Asn 330 Met Val Leu Thr Glu Thr Gly Ala Ala Lys Ala Val Ser Lys Ala Leu Pro Glu Leu Glu Gly Lys Leu Thr Gly Asn Ala Ile Arg Val Pro Thr Pro Asp Val Ser Met Ala Val Leu Asn Leu Thr Leu Asn Thr Glu Val 375 Asp Arg Asp Glu Val Asn Glu Phe Leu Arg Arg Val Ser Leu His Ser 385 395 Asp Leu Arg Gln Gln Ile Asp Trp Ile Arg Ser Pro Glu Val Val Ser 410

Thr Asp Phe Val Gly Thr Thr His Ala Gly Ile Val Asp Gly Leu Ala 425

Thr Ile Ala Thr Gly Arg His Leu Val Leu Tyr Val Trp Tyr Asp Asn 435

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				gcc Ala												547
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aac Asn	atc Ile	gtg Val	tac Tyr 265	ggc Gly	atc Ile	aac Asn	cac His	acc Thr 270	gac Asp	atc Ile	acc Thr	gca Ala	gat Asp 275	gat Asp	cag Gln	931
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					ctg Leu											787
					gtt Val 235											835
					tct Ser											883
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- Asp Ser Ile Met Gly Arg Leu Gly Gln Glu Val Glu Tyr Asp Asp Asp 50 55 60
- Ser Ile Thr Val Gly Gly Lys Arg Ile Ala Val Tyr Ala Glu Arg Asp
 65 70 75 80
- Pro Lys Asn Leu Asp Trp Ala Ala His Asn Val Asp Ile Val Ile Glu 85 90 95
- Ser Thr Gly Phe Phe Thr Asp Ala Asn Ala Ala Lys Ala His Ile Glu 100 105 110
- Ala Gly Ala Lys Lys Val Ile Ile Ser Ala Pro Ala Ser Asn Glu Asp 115 120 125
- Ala Thr Phe Val Tyr Gly Val Asn His Glu Ser Tyr Asp Pro Glu Asn 130 135 140
- His Asn Val Ile Ser Gly Ala Ser Cys Thr Thr Asn Cys Leu Ala Pro 145 150 155 160
- Met Ala Lys Val Leu Asn Asp Lys Phe Gly Ile Glu Asn Gly Leu Met 165 170 175
- Thr Thr Val His Ala Tyr Thr Gly Asp Gln Arg Leu His Asp Ala Pro 180 185 190
- His Arg Asp Leu Arg Arg Ala Arg Ala Ala Ala Val Asn Ile Val Pro 195 200 205
- Thr Ser Thr Gly Ala Ala Lys Ala Val Ala Leu Val Leu Pro Glu Leu 210 215 220
- Lys Gly Lys Leu Asp Gly Tyr Ala Leu Arg Val Pro Val Ile Thr Gly
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- Ser Ala Thr Asp Leu Thr Phe Asn Thr Lys Ser Glu Val Thr Val Glu 245 250 255
- Ser Ile Asn Ala Ala Ile Lys Glu Ala Ala Val Gly Glu Phe Gly Glu 260 265 270
- Thr Leu Ala Tyr Ser Glu Glu Pro Leu Val Ser Thr Asp Ile Val His 275 280 285
- Asp Ser His Gly Ser Ile Phe Asp Ala Gly Leu Thr Lys Val Ser Gly 290 295 300
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						Ser				acc Thr 320						1075
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Lys Ala Leu Ser Glu Gly Gly Ala Lys Val Ile Val Met Ala His Leu 50 55 60

Gly Arg Pro Lys Gly Glu Val Asn Glu Lys Tyr Ser Leu Ala Pro Val 65 70 75 80

Ala Glu Ala Leu Ser Asp Glu Leu Gly Gln Tyr Val Ala Leu Ala Ala 85 90 95

Asp Val Val Gly Glu Asp Ala His Glu Arg Ala Asn Gly Leu Thr Glu
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Gly Asp Ile Leu Leu Glu Asn Val Arg Phe Asp Pro Arg Glu Thr 115 120 125

Ser Lys Asp Glu Ala Glu Arg Thr Ala Phe Ala Gln Glu Leu Ala Ala 130 135 140

Leu Ala Ala Asp Asn Gly Ala Phe Val Ser Asp Gly Phe Gly Val Val 145 150 155 160

His Arg Ala Gln Thr Ser Val Tyr Asp Ile Ala Lys Leu Leu Pro His 165 170 175

Tyr Ala Gly Gly Leu Val Glu Thr Glu Ile Ser Val Leu Glu Lys Ile 180 185 190

Ala Glu Ser Pro Glu Ala Pro Tyr Val Val Leu Gly Gly Ser Lys 195 200 205

Val Ser Asp Lys Ile Gly Val Ile Glu Ala Leu Ala Ala Lys Ala Asp 210 215 220

Lys Ile Ile Val Gly Gly Met Cys Tyr Thr Phe Leu Ala Ala Gln 225 230 235 240

Gly His Asn Val Gln Gln Ser Leu Leu Gln Glu Glu Met Lys Ala Thr 245 250 255

Cys Thr Asp Leu Leu Ala Arg Phe Gly Asp Lys Ile Val Leu Pro Val 260 265 270

Asp Leu Val Ala Ala Ser Glu Phe Asn Lys Asp Ala Glu Lys Gln Ile 275 280 285

Val Asp Leu Asp Ser Ile Pro Glu Gly Trp Met Ser Leu Asp Ile Gly Pro Glu Ser Val Lys Asn Phe Gly Glu Val Leu Ser Thr Ala Lys Thr 310 Ile Phe Trp Asn Gly Pro Met Gly Val Phe Glu Phe Ala Ala Phe Ser 325 330 Glu Gly Thr Arg Gly Ile Ala Gln Ala Ile Ile Asp Ala Thr Ala Gly Asn Asp Ala Phe Ser Val Val Gly Gly Asp Ser Ala Ala Ser Val 360 Arg Val Leu Gly Leu Asn Glu Asp Gly Phe Ser His Ile Ser Thr Gly 370 375 Gly Gly Ala Ser Leu Glu Tyr Leu Glu Gly Lys Glu Leu Pro Gly Val 390 395 Ala Ile Leu Ala Gln 405 <210> 71 <211> 1398 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1375) <223> RXA00235 <400> 71 cgaaacaaga ttcgtgcaac aattgggtgt agacgtgatt gaagacattt gatcacgtga 60 ataattctag ttagctccca agttggcata ggaggccaca gtg gct gaa atc atg Val Ala Glu Ile Met cac gta ttc gct cgc gaa att ctc gac tcc cgc ggt aac cca acc gtc 163 His Val Phe Ala Arg Glu Ile Leu Asp Ser Arg Gly Asn Pro Thr Val 3.5 gag gca gag gtt ttc ctg gat gac ggt tcc cac ggt gtc gca ggt gtt 211 Glu Ala Glu Val Phe Leu Asp Asp Gly Ser His Gly Val Ala Gly Val cea tee gge gea tee ace gge gte cae gag get cat gag etg egt gae 259 Pro Ser Gly Ala Ser Thr Gly Val His Glu Ala His Glu Leu Arg Asp 40 45 50 ggt ggc gat cgc tac ctg ggc aag ggc gtt ttg aag gca gtt gaa aac 307 Gly Gly Asp Arg Tyr Leu Gly Lys Gly Val Leu Lys Ala Val Glu Asn gtc aac gaa gaa atc ggc gac gag ctc gct ggc cta gag gct gac gat 355 Val Asn Glu Glu Ile Gly Asp Glu Leu Ala Gly Leu Glu Ala Asp Asp 70 75 80

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			gca Ala							451
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			cac His							547
			gct Ala 155							595
			gca Ala							643
			gca Ala							691
			gat Asp							739
			gac Asp							787
			gac Asp 235							835
	-	_	ggc Gly			_	 	_		883
			aac Asn							931
			gac Asp							979
			acc Thr							1027
			aac Asn 315							1075

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gca Ala 390	cgt Arg	tcc Ser	gac Asp	cgt Arg	gtc Val 395	gca Ala	aag Lys	tac Tyr	aac Asn	cag Gln 400	ctt Leu	ctc Leu	cgc Arg	atc Ile	gag Glu 405	1315	
cag Gln	ctg Leu	ctt Leu	ggc Gly	gac Asp 410	gcc Ala	ggc Gly	gtc Val	tac Tyr	gca Ala 415	ggt Gly	cgc Arg	agc Ser	gca Ala	ttc Phe 420	cca Pro	1363	;
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Met Met Asn Ile Ile Thr Gly Gly Ala His Ala Asp Ser Gly Val Asp Val Gln Glu Phe Met Ile Ala Pro Ile Gly Ala Glu Thr Phe Ser Glu 170 165 Ala Leu Arg Asn Gly Ala Glu Val Tyr His Ala Leu Lys Ser Val Ile 185 Lys Glu Lys Gly Leu Ser Thr Gly Leu Gly Asp Glu Gly Gly Phe Ala Pro Ser Val Gly Ser Thr Arg Glu Ala Leu Asp Leu Ile Val Glu Ala Ile Glu Lys Ala Gly Phe Thr Pro Gly Lys Asp Ile Ala Leu Ala Leu Asp Val Ala Ser Ser Glu Phe Phe Lys Asp Gly Thr Tyr His Phe Glu 245 Gly Gly Gln His Ser Ala Ala Glu Met Ala Asn Val Tyr Ala Glu Leu 265 Val Asp Ala Tyr Pro Ile Val Ser Ile Glu Asp Pro Leu Gln Glu Asp Asp Trp Glu Gly Tyr Thr Asn Leu Thr Ala Thr Ile Gly Asp Lys Val Gln Ile Val Gly Asp Asp Phe Phe Val Thr Asn Pro Glu Arg Leu Lys 315 310 Glu Gly Ile Ala Lys Lys Ala Ala Asn Ser Ile Leu Val Lys Val Asn Gln Ile Gly Thr Leu Thr Glu Thr Phe Asp Ala Val Asp Met Ala His 345 Arg Ala Gly Tyr Thr Ser Met Met Ser His Arg Ser Gly Glu Thr Glu Asp Thr Thr Ile Ala Asp Leu Ala Val Ala Leu Asn Cys Gly Gln Ile 375 Lys Thr Gly Ala Pro Ala Arg Ser Asp Arg Val Ala Lys Tyr Asn Gln 385 Leu Leu Arg Ile Glu Gln Leu Leu Gly Asp Ala Gly Val Tyr Ala Gly Arg Ser Ala Phe Pro Arg Phe Gln Gly 420

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180

691

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Ala Leu Lys Leu Gly Val Asp Phe Ile Ala Leu Ser Phe Val Arg Ser

cca gca gat gct gaa ctc gtt cac aag atc atg gac gaa gaa ggt cgt

190

185

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	gtt Val 215															787
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	gca Ala															931
	cag Gln															979
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	ctt Leu															1075
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	gca Ala															1219
	acc Thr 375															1267
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cct Pro	gag Glu	tac Tyr	aac Asn	aag Lys	ggt Gly	gac Asp	atg Met	atg Met	gtt Val	gtt Val	gtt Val	gca Ala	ggt Gly	tcc Ser	cct Pro	1459

440 445 450

cct ggt gtt acc ggt aac acc aac atg att cac gtc cac ctt ctt ggt
Pro Gly Val Thr Gly Asn Thr Asn Met Ile His Val His Leu Leu Gly
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Val Ala Arg Leu Asn Phe Ser His Gly Asp His Pro Asp His Glu Gln 35 40 45

Asn Tyr Lys Trp Val Arg Glu Ala Ala Glu Lys Thr Gly Arg Ala Val
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Gly Ile Leu Ala Asp Leu Gln Gly Pro Lys Ile Arg Leu Gly Arg Phe
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Thr Asp Gly Ala Thr Val Trp Glu Asn Gly Glu Thr Ile Arg Ile Thr 85 90 .95

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Asn Leu Ala Lys Asp Ala Lys Pro Gly Asp Arg Leu Leu Val Asp Asp 115 120 125

Gly Lys Val Gly Leu Val Cys Val Ser Val Glu Gly Asn Asp Val Ile 130 135 140

Cys Glu Val Val Glu Gly Gly Pro Val Ser Asn Asn Lys Gly Val Ser 145 150 155 160

Leu Pro Gly Met Asp Ile Ser Val Pro Ala Leu Ser Glu Lys Asp Ile 165 170 175

Arg Asp Leu Arg Phe Ala Leu Lys Leu Gly Val Asp Phe Ile Ala Leu 180 185 190

Ser Phe Val Arg Ser Pro Ala Asp Ala Glu Leu Val His Lys Ile Met 195 200 205

Asp Glu Glu Gly Arg Arg Val Pro Val Ile Ala Lys Leu Glu Lys Pro 210 215 220

Glu Ala Val Thr Ser Leu Glu Pro Ile Val Leu Ala Phe Asp Ala Val 225 230 235 240

Met Val Ala Arg Gly Asp Leu Gly Val Glu Val Pro Leu Glu Glu Val Pro Leu Val Gln Lys Arg Ala Ile Gln Ile Ala Arg Glu Asn Ala Lys 265 Pro Val Ile Val Ala Thr Gln Met Leu Asp Ser Met Ile Glu Asn Ser 275 280 Arg Pro Thr Arg Ala Glu Ala Ser Asp Val Ala Asn Ala Val Leu Asp Gly Ala Asp Ala Val Met Leu Ser Gly Glu Thr Ser Val Gly Lys Asp 310 Pro His Asn Val Val Arg Thr Met Ser Arg Ile Val Arg Phe Ala Glu 330 Thr Asp Gly Arg Val Pro Asp Leu Thr His Ile Pro Arg Thr Lys Arg 340 Gly Val Ile Ser Tyr Ser Ala Arg Asp Ile Ala Glu Arg Leu Asn Ala 360 Arg Ala Leu Val Ala Phe Thr Thr Ser Gly Asp Thr Ala Lys Arg Val 370 380 Ala Arg Leu His Ser His Leu Pro Leu Leu Val Phe Thr Pro Asn Glu Ala Val Arg Ser Glu Leu Ala Leu Thr Trp Gly Ala Thr Thr Phe Leu 410 Cys Pro Pro Val Ser Asp Thr Asp Asp Met Met Arg Glu Val Asp Arg 425 Ala Leu Leu Ala Met Pro Glu Tyr Asn Lys Gly Asp Met Met Val Val 440 Val Ala Gly Ser Pro Pro Gly Val Thr Gly Asn Thr Asn Met Ile His 455 450 Val His Leu Leu Gly Asp Asp Thr Arg Ile Ala Lys Leu 470 <210> 75 <211> 1980 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1957) <223> RXN02675 <400> 75 aagtgtttca ttggaacact tgcgctgcca actttttggt ttacgggcac aatgaaactg 60

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cag gac att ctc cag gag atc aag act gaa ctc gac gag tta att cta 163 Gln Asp Ile Leu Gln Glu Ile Lys Thr Glu Leu Asp Glu Leu Ile Leu 10 15 20

gaa ctt gat gag gtg aca caa act cac agc gag gcc atc ggg cag gtc 211 Glu Leu Asp Glu Val Thr Gln Thr His Ser Glu Ala Ile Gly Gln Val 25 30 35

tcc cca acc cat tac gtt ggt gcc cgc aac ctc atg cat tac gcg cat 259 Ser Pro Thr His Tyr Val Gly Ala Arg Asn Leu Met His Tyr Ala His 40 45 50

ctt cgc acc aaa gac ctc cgt ggc ctg cag caa cgc ctc tcc tct gtg 307 Leu Arg Thr Lys Asp Leu Arg Gly Leu Gln Gln Arg Leu Ser Ser Val

gga gct acc cgc ttg act acc acc gaa cca gca gtg cag gcc cgc ctc 355
Gly Ala Thr Arg Leu Thr Thr Glu Pro Ala Val Gln Ala Arg Leu
70 75 80 85

aag gcc gcc cgc aat gtt atc gga gct ttc gca ggt gaa ggc cca ctt 403 Lys Ala Ala Arg Asn Val Ile Gly Ala Phe Ala Gly Glu Gly Pro Leu

tat cca ccc tca gat gtc gtc gat gcc ttc gaa gat gcc gat gag att 451
Tyr Pro Pro Ser Asp Val Val Asp Ala Phe Glu Asp Ala Asp Glu Ile
105 110 115

ctc gac gag cac gcc gaa att ctc ctt ggc gaa ccc cta ccg gat act 499 Leu Asp Glu His Ala Glu Ile Leu Leu Gly Glu Pro Leu Pro Asp Thr 120 125 130

cca tcc tgc atc atg gtc acc ctg ccc acc gaa gcc gcc acc gac att 547
Pro Ser Cys Ile Met Val Thr Leu Pro Thr Glu Ala Ala Thr Asp Ile
135 140 145

gaa ctt gtc cgt ggc ttc gcc aaa agc ggc atg aat cta gct cgc atc 595 Glu Leu Val Arg Gly Phe Ala Lys Ser Gly Met Asn Leu Ala Arg Ile 150 165

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gac ctc gcc gga cca aaa gta cgc acc ggc gaa atc gcc cca ggc gca 739
Asp Leu Ala Gly Pro Lys Val Arg Thr Gly Glu Ile Ala Pro Gly Ala
200 205 210

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acg ccc gca aaa ctg tgg atc acc gcc cac ggc tcc gaa cca gtc cca 835 Thr Pro Ala Lys Leu Trp Ile Thr Ala His Gly Ser Glu Pro Val Pro

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cca gaa tgg tto Pro Glu Trp Pho 26	e Asp Lys Leu	gaa atc ggc Glu Ile Gly 270	agc gtc atc aac gt Ser Val Ile Asn Va 275	c cca 931 l Pro
gac acc cgc gga Asp Thr Arg Gly 280	a toc ogo oga y Ser Arg Arg	gca ttc acc Ala Phe Thr 285	gtg acc agg gtt tt Val Thr Arg Val Ph 290	t gat 979 e Asp
ggc gcg gtc ctc Gly Ala Val Len 295	c gcc gaa ggc ı Ala Glu Gly 300	cca caa aaa Pro Gln Lys	gcc tac atc tcc aa Ala Tyr Ile Ser As 305	c ggc 1027 n Gly
acc ctc ctg gad Thr Leu Leu Glu 310	a cac aac tac ı His Asn Tyr 315	gac cgc tcc Asp Arg Ser	cgg gtc tac ggc at Arg Val Tyr Gly Il 320	c ccc 1075 e Pro 325
gcc gta gtt cac Ala Val Val Gl	g cgc atc aac n Arg Ile Asn 330	ctc aaa gtc Leu Lys Val 335	ggc gac cgc ctc at Gly Asp Arg Leu Il 34	e Leu
acc gac gaa gaa Thr Asp Glu Gla 34	u Leu Thr Tyr	gat cca tcc Asp Pro Ser 350	ctc gga tcc ggc cg Leu Gly Ser Gly Ar 355	c aca 1171 g Thr
cca cgc atc ago Pro Arg Ile Se 360	c tgc acc ctt r Cys Thr Leu	cca caa gca Pro Gln Ala 365	gtc gat gca att aa Val Asp Ala Ile Ly 370	a gtc 1219 s Val
ggg cac cgc gtc Gly His Arg Va 375	g ctt ttc gac l Leu Phe Asp 380	gac gga gcc Asp Gly Ala	atc gcc gca gtc tg Ile Ala Ala Val Cy 385	c atc 1267 s Ile
gac aag acc tc Asp Lys Thr Se 390	r Thr Ala Asp	Gly His Asn	gac gta gaa ttg ga Asp Val Glu Leu Gl 400	a gtc 1315 u Val 405
acc cac gcc cg Thr His Ala Arc	c cca caa ggc g Pro Gln Gly 410	gta aac ctg Val Asn Leu 415	gcc gca tac aag gg Ala Ala Tyr Lys Gl 42	y Ile
aac ctc cca gad Asn Leu Pro Asp 42	p Ser Glu Leu	cca ctc cca Pro Leu Pro 430	agc ctc act gaa ga Ser Leu Thr Glu Gl 435	a gac 1411 u Asp
ctc caa cac ctc Leu Gln His Lev 440	g cgc ttt gtc u Arg Phe Val	gtc aaa tac Val Lys Tyr 445	gcc gac atc gca gc Ala Asp Ile Ala Al 450	c atc 1459 a Ile
tcc ttc atc cga Ser Phe Ile Arc 455	a aaç gtc gcc g Asn Val Ala 460	gac gtg gaa Asp Val Glu	tac ctc ctc caa go Tyr Leu Leu Gln Al 465	a ctc 1507 a Leu
gcc gac atc gga Ala Asp Ile Gl 470	a gat cca gta y Asp Pro Val 475	gcc gtc gaa Ala Val Glu	cgc ctt ggc ctc gt Arg Leu Gly Leu Va 480	c ctt 1555 l Leu 485

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										atg Met						1651
										gca Ala						1699
										cca Pro						1747
										ctc Leu 560						1795
										gaa Glu						1843
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Ala	Ile	Gly 35	Gln	Val	Ser	Pro	Thr 40	His	Tyr	Val	Gly	Ala 45	Arg	Asn	Leu	
Met	His 50	Tyr	Ala	His	Leu	Arg 55	Thr	Lys	Asp	Leu	Arg 60	Gly	Leu	Gln	Gln	
Arg 65	Leu	Ser	Ser	Val	Gly 70	Ala	Thr	Arg	Leu	Thr 75	Thr	Thr	Glu	Pro	Ala 80	
Val	Gln	Ala	Arg	Leu 85	Lys	Ala	Ala	Arg	Asn 90	Val	Ile	Gly	Ala	Phe 95	Ala	

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Ala Tyr Lys Gly Ile Asn Leu Pro Asp Ser Glu Leu Pro Leu Pro Ser Leu Thr Glu Glu Asp Leu Gln His Leu Arg Phe Val Val Lys Tyr Ala 440 Asp Ile Ala Ala Ile Ser Phe Ile Arg Asn Val Ala Asp Val Glu Tyr Leu Leu Gln Ala Leu Ala Asp Ile Gly Asp Pro Val Ala Val Glu Arg 470 475 Leu Gly Leu Val Leu Lys Ile Glu Thr Ile Pro Gly Tyr Glu Gly Leu 485 Ala Gln Ile Leu Leu Thr Gly Met Arg His Glu Asn Phe Gly Ile Met 500 Ile Ala Arg Gly Asp Leu Ala Val Glu Leu Gly Phe Asp Arg Met Ala 520 Glu Val Pro Gln Leu Ile Met Ala Leu Ala Glu Ala Ala His Val Pro 535 Thr Ile Leu Ala Thr Gln Val Leu Glu Asn Met Ala Lys Asn Gly Leu 550 555 Pro Ser Arg Ala Glu Ile Thr Asp Ala Ala Met Ala Leu Arg Ala Glu 570 Cys Val Met Leu Asn Lys Gly Pro His Ile Asn Asp Ala Ile Lys Val 580 585 Leu Thr Glu Met Ser Arg Lys Leu Gly Ala Ser Gln Arg Lys Ser Arg 600 Leu Leu Arg Lys Val Lys Ser Trp Glu Glu 610 <210> 77 <211> 386 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (1)..(363) <223> FRXA02675 <400> 77 ate ete atg ace gge atg ege cae gaa aac tte gge ate atg ate gee Ile Leu Met Thr Gly Met Arg His Glu Asn Phe Gly Ile Met Ile Ala 1 cgc gga gac ctc gcc gtc gaa ctc ggc ttc gac cgc atq qca qaa qtc 96 Arg Gly Asp Leu Ala Val Glu Leu Gly Phe Asp Arg Met Ala Glu Val 20 25 ccc caa ctg atc atg gcc ctt gca gaa gcc gcc cac gtc cca acc atc

Pro Gln Leu Ile Met Ala Leu Ala Glu Ala Ala His Val Pro Thr Ile

35 40 45 192 ttq gcc acc caa gtc ctg gaa aac atg gcc aaa aac gga ctc cca tct Leu Ala Thr Gln Val Leu Glu Asn Met Ala Lys Asn Gly Leu Pro Ser 55 cgc gca gaa atc acc gac gca gca atg gca ctt cgc gct gaa tgc gtc 240 Arg Ala Glu Ile Thr Asp Ala Ala Met Ala Leu Arg Ala Glu Cys Val 70 75 288 atg ctg aac aag gga cca cac atc aac gac gcc atc aag gtc ctc acc Met Leu Asn Lys Gly Pro His Ile Asn Asp Ala Ile Lys Val Leu Thr 90 gaa atg agc cgc aaa ctt ggt gca tcc caa cga aag agt agg ctg ctg 336 Glu Met Ser Arg Lys Leu Gly Ala Ser Gln Arg Lys Ser Arg Leu Leu 105 100 383 ctg cqc aag gtg aag agc tgg gaa gag taactcacaa aggcgattgg Leu Arg Lys Val Lys Ser Trp Glu Glu 115 386 cgt <210> 78 <211> 121 <212> PRT <213> Corynebacterium glutamicum <400> 78 Ile Leu Met Thr Gly Met Arg His Glu Asn Phe Gly Ile Met Ile Ala Arg Gly Asp Leu Ala Val Glu Leu Gly Phe Asp Arg Met Ala Glu Val Pro Gln Leu Ile Met Ala Leu Ala Glu Ala Ala His Val Pro Thr Ile Leu Ala Thr Gln Val Leu Glu Asn Met Ala Lys Asn Gly Leu Pro Ser 55 50 Arg Ala Glu Ile Thr Asp Ala Ala Met Ala Leu Arg Ala Glu Cys Val 70 Met Leu Asn Lys Gly Pro His Ile Asn Asp Ala Ile Lys Val Leu Thr Glu Met Ser Arg Lys Leu Gly Ala Ser Gln Arg Lys Ser Arg Leu Leu 105

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Asp	Leu	Ala	Gly	Pro	Lys	Val	Arg	Thr	Gly	Glu	Ile	Ala	Pro	Gly	Ala	
•		200	-		-		205		-			210		-		
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				ctg Leu											cca Pro 245	835
				ctg Leu 250												883
				gac Asp												931
				tcc Ser		Arg										979
		-		gcc Ala	-					-						1027
				cac His												1075
				cgc Arg 330												1123
				ctc Leu												1171
				tgc Cys												1219
				ctt Leu												1267
				act Thr												1315
				cca Pro 410												1363
				tcc Ser												1411
ctc Leu	caa Gln	cac His	ctg Leu	cgc Arg	ttt Phe	gtc Val	gtc Val	aaa Lys	tac Tyr	gcc Ala	gac Asp	atc Ile	gca Ala	gcc Ala	atc Ile	1459

440 445 450

tcc ttc atc cga aac gtc gcc gac gtg gaa tac ctc ctc caa gca ctc

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455

460

465

gcc gac atc gga gat

Ala Asp Ile Gly Asp
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Ala Ile Gly Gln Val Ser Pro Thr His Tyr Val Gly Ala Arg Asn Leu $35 \hspace{1cm} 40 \hspace{1cm} 45$

Met His Tyr Ala His Leu Arg Thr Lys Asp Leu Arg Gly Leu Gln Gln 50 55 60

Arg Leu Ser Ser Val Gly Ala Thr Arg Leu Thr Thr Thr Glu Pro Ala 65 70 75 80

Val Gln Ala Arg Leu Lys Ala Ala Arg Asn Val Ile Gly Ala Phe Ala 85 90 95

Gly Glu Gly Pro Leu Tyr Pro Pro Ser Asp Val Val Asp Ala Phe Glu 100 105 110

Asp Ala Asp Glu Ile Leu Asp Glu His Ala Glu Ile Leu Gly Glu
115 120 125

Pro Leu Pro Asp Thr Pro Ser Cys Ile Met Val Thr Leu Pro Thr Glu 130 135 140

Ala Ala Thr Asp Ile Glu Leu Val Arg Gly Phe Ala Lys Ser Gly Met 145 150 155 160

Asn Leu Ala Arg Ile Asn Cys Ala His Asp Asp Glu Thr Val Trp Lys 165 170 175

Gln Met Ile Asp Asn Val His Thr Val Ala Glu Glu Val Gly Arg Glu 180 185 190

Ile Arg Val Ser Met Asp Leu Ala Gly Pro Lys Val Arg Thr Gly Glu
195 200 205

Ile Ala Pro Gly Ala Glu Val Gly Arg Ala Arg Val Thr Arg Asp Glu 210 215 220

Thr Gly Lys Val Leu Thr Pro Ala Lys Leu Trp Ile Thr Ala His Gly 225 230 235 240

Ser Glu Pro Val Pro Ala Pro Glu Ser Leu Pro Gly Arg Pro Ala Leu 245 250 255

Pro Ile Glu Val Thr Pro Glu Trp Phe Asp Lys Leu Glu Ile Gly Ser 260 265 270

Val Ile Asn Val Pro Asp Thr Arg Gly Ser Arg Arg Ala Phe Thr Val 275 280 285

Thr Arg Val Phe Asp Gly Ala Val Leu Ala Glu Gly Pro Gln Lys Ala 290 295 300

Tyr Ile Ser Asn Gly Thr Leu Leu Glu His Asn Tyr Asp Arg Ser Arg 305 310 315 320

Val Tyr Gly Ile Pro Ala Val Val Gln Arg Ile Asn Leu Lys Val Gly 325 330 335

Asp Arg Leu Ile Leu Thr Asp Glu Glu Leu Thr Tyr Asp Pro Ser Leu 340 345 350

Gly Ser Gly Arg Thr Pro Arg Ile Ser Cys Thr Leu Pro Gln Ala Val 355 360 365

Asp Ala Ile Lys Val Gly His Arg Val Leu Phe Asp Asp Gly Ala Ile 370 375 380

Ala Ala Val Cys Ile Asp Lys Thr Ser Thr Ala Asp Gly His Asn Asp 385 390 395 400

Val Glu Leu Glu Val Thr His Ala Arg Pro Gln Gly Val Asn Leu Ala 405 410 415

Ala Tyr Lys Gly Ile Asn Leu Pro Asp Ser Glu Leu Pro Leu Pro Ser 420 425 430

Leu Thr Glu Glu Asp Leu Gln His Leu Arg Phe Val Val Lys Tyr Ala 435 440 445

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Met Ala Asn Lys Ser ttc ccc aag ccc tcc gat ctt cca gtg ccc aag ggc gct gaa ggt tgg 163 Phe Pro Lys Pro Ser Asp Leu Pro Val Pro Lys Gly Ala Glu Gly Trp gaa gat ctg tac ccg tac tac ctc gtt ttc caa gac aag ctc atg gat 211 Glu Asp Leu Tyr Pro Tyr Tyr Leu Val Phe Gln Asp Lys Leu Met Asp caa gag aat gag aaa ttc tgg ttc tgc gat tca cag cac tgg cca act 259 Gln Glu Asn Glu Lys Phe Trp Phe Cys Asp Ser Gln His Trp Pro Thr 45 gtg ttc aag cct ttt gaa act atc ggt ggt gaa ttc gct gta aag tgc 307 Val Phe Lys Pro Phe Glu Thr Ile Gly Gly Glu Phe Ala Val Lys Cys ctc ggc caa tac aac gct cgg cat ttg atg atc ccg aat gcc aat ggc 355 Leu Gly Gln Tyr Asn Ala Arg His Leu Met Ile Pro Asn Ala Asn Gly 80 atc gag ttc cgc gtg cat ctg gga tac ctc tat atg tcc cct att cca 403 Ile Glu Phe Arg Val His Leu Gly Tyr Leu Tyr Met Ser Pro Ile Pro gtg cct gaa gat cag att gcg gaa cgc gtc ccc atg ttc cag gaa cgc 451 Val Pro Glu Asp Gln Ile Ala Glu Arg Val Pro Met Phe Gln Glu Arg 105 110 atc acg cac tac ttc caa aac tgg gag cca atg ctg gca aat tgg aag Ile Thr His Tyr Phe Gln Asn Trp Glu Pro Met Leu Ala Asn Trp Lys 120 125 gag cga gta tta gga acc atc aat gag ctg gaa tct cta gaa ttc aag 547 Glu Arg Val Leu Gly Thr Ile Asn Glu Leu Glu Ser Leu Glu Phe Lys 135 cca ctg cct gac tac gtg cct atc gat gat att gtc tcc gga aaa gcc Pro Leu Pro Asp Tyr Val Pro Ile Asp Asp Ile Val Ser Gly Lys Ala 150 155 aaa gac ggc acc gaa gta ctc atg gaa aac ttc gat cgg ctc att cag Lys Asp Gly Thr Glu Val Leu Met Glu Asn Phe Asp Arg Leu Ile Gln 170 ctc gcc tac caa aac tgg caa tac cac ttt gag ttc ctc aac ttg ggt Leu Ala Tyr Gln Asn Trp Gln Tyr His Phe Glu Phe Leu Asn Leu Gly 185 190 tac atc gct tac cta gat ttc ttc aat ttc tgc aag gaa gtc ttc cca 739 Tyr Ile Ala Tyr Leu Asp Phe Phe Asn Phe Cys Lys Glu Val Phe Pro 200 gat atc cct gat caa tca att tcg atg atg gtt cag ggc gtg gat atg 787 Asp Ile Pro Asp Gln Ser Ile Ser Met Met Val Gln Gly Val Asp Met 215 220 gag ctg ttc cgc ccc gat gat gaa cta aag att ctg gca cag cta gcg 835 Glu Leu Phe Arg Pro Asp Asp Glu Leu Lys Ile Leu Ala Gln Leu Ala

230					235					240					245	
													gat Asp			883
													aca Thr 275			931
													acc Thr			979
													ctg Leu			1027
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	Arg	Glu	Leu	Ser	Arg	Thr	Leu	Gln	Gly	Tyr	Gly	Phe	tgg Trp	Glu		1315
													gat Asp			1363
													ggt Gly 435			1411
													gca Ala			1459
.acc Thr													act Thr			1507
													atc Ile			1555

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acc Thr	ctt Leu	aaa Lys	ggc Gly 505	atg Met	gct Ala	gca Ala	tcc Ser	cct Pro 510	ggt Gly	gtg Val	gtg Val	gaa Glu	ggc Gly 515	tac Tyr	gct Ala	1651
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								cct Pro								1747
								att Ile								1795
								ttg Leu								1843
								ggc Gly 590								1891
acc Thr	aag Lys	ggc Gly 600	aag Lys	gtt Val	gtc Val	att Ile	gtt Val 605	gat Asp	cca Pro	gat Asp	gcg Ala	cca Pro 610	cgc Arg	atc Ile	gaa Glu	1939
								cac His								1987
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Gly	Ala	Glu	Gly 20	Trp	Glu	Asp	Leu	Tyr 25	Pro	Tyr	Tyr	Leu	Val 30	Phe	Gln	
Asp	Lys	Leu 35	Met	Asp	Gln	Glu	Asn 40	Glu	Lys	Phe	Trp	Phe 45	Cys	Asp	Ser	

122

Gln His Trp Pro Thr Val Phe Lys Pro Phe Glu Thr Ile Gly Gly Glu

Phe Ala Val Lys Cys Leu Gly Gln Tyr Asn Ala Arg His Leu Met Ile

55

50

70 75 65 Pro Asn Ala Asn Gly Ile Glu Phe Arg Val His Leu Gly Tyr Leu Tyr 85 Met Ser Pro Ile Pro Val Pro Glu Asp Gln Ile Ala Glu Arg Val Pro 105 Met Phe Gln Glu Arg Ile Thr His Tyr Phe Gln Asn Trp Glu Pro Met 120 Leu Ala Asn Trp Lys Glu Arg Val Leu Gly Thr Ile Asn Glu Leu Glu Ser Leu Glu Phe Lys Pro Leu Pro Asp Tyr Val Pro Ile Asp Asp Ile 155 150 Val Ser Gly Lys Ala Lys Asp Gly Thr Glu Val Leu Met Glu Asn Phe Asp Arg Leu Ile Gln Leu Ala Tyr Gln Asn Trp Gln Tyr His Phe Glu 185 Phe Leu Asn Leu Gly Tyr Ile Ala Tyr Leu Asp Phe Phe Asn Phe Cys 200 Lys Glu Val Phe Pro Asp Ile Pro Asp Gln Ser Ile Ser Met Met Val 215 Gln Gly Val Asp Met Glu Leu Phe Arg Pro Asp Asp Glu Leu Lys Ile 235 Leu Ala Gln Leu Ala Val Asp Leu Gly Leu Gln Thr His Phe Ala Asn Pro Asp Asp Pro Gln Ala Thr Leu Ala Ala Ile Ala Lys Ala Glu Gly 265 Gly Ala Thr Trp Ile Ala Arg Trp Glu Glu Ala Gln Asp Pro Trp Phe Asn Phe Thr Val Gly Asn Gly Phe Tyr Gly His Asp Lys Tyr Trp Ile 295 Glu His Leu Glu Leu Pro Leu Gly Tyr Ile Ala Asp Tyr Ile Arg Arg 305 Leu Asp Glu Gly Gln Thr Ile Ser Arg Pro Lys Asp Glu Leu Ile Ala 330 Glu Lys Glu Arg Val Val Glu Glu Tyr Arg Asp Leu Leu Asp Gly Glu 340 Gln Leu Ala Gln Phe Asp Ala Lys Cys Gly Leu Ala Ala Thr Ala Tyr Pro Tyr Val Glu Asn His Asn Phe Tyr Ile Glu His Trp Thr Met Ser 370 375 Val Phe Trp Arg Lys Val Arg Glu Leu Ser Arg Thr Leu Gln Gly Tyr 385 390

Gly Phe Trp Glu Asn Glu Asp Asp Met Leu Tyr Leu Asn Arg Thr Glu 405 Val Arg Asp Val Leu Phe Asp Leu Ala Thr Ala Trp Gly Val Gly Ala 425 Pro Gly Gly Pro Ile Gly Thr Ile Ile Trp Pro Glu Glu Ile Glu Arg 440 Arg Lys Ala Ile Val Thr Ala Leu Lys Thr Ala Arg Pro Ala Pro Ala Leu Asn Thr Pro Pro Glu Ser Ile Thr Glu Pro Phe Thr Arg Met Leu 475 Trp Gly Ile Thr Thr Glu Gln Val Gln Ser Trp Leu Gly Asn Asp Glu 490 Asp Ala Glu Glu Gly Thr Leu Lys Gly Met Ala Ala Ser Pro Gly Val Val Glu Gly Tyr Ala Arg Val Ile Leu Ser Ala Asp Asp Leu Ser Glu 520 Ile Gln Gln Asp Glu Ile Leu Val Ala Pro Val Thr Ala Pro Ser Trp 535 Gly Pro Ile Phe Gly Lys Ile Lys Ala Thr Val Thr Asp Ile Gly Gly 550 Met Met Ser His Ala Ala Ile Val Cys Arg Glu Tyr Gly Leu Pro Ala 570 Val Thr Gly Thr Gly Ala Ala Ser Thr Thr Ile Lys Thr Gly Asp Tyr Leu Lys Val Asp Gly Thr Lys Gly Lys Val Val Ile Val Asp Pro Asp 600 Ala Pro Arg Ile Glu Gly Pro Gly Ala His Ser His Ala His Ser Val Ala Ala His Gly Val Asp Thr His Ala 625 630 <210> 83 <211> 1215 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1192) <223> RXA00683

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Met Thr Asn Ser Leu aac atc ccg ttt gtc cag cgc ttc gat gaa ggc ctg gat cct gtt cta 163 Asn Ile Pro Phe Val Gln Arg Phe Asp Glu Gly Leu Asp Pro Val Leu 10 gaa qta ctc qqt qqc aaq qqc qct tca cta qtc acc atg aca gat qct 211 Glu Val Leu Gly Gly Lys Gly Ala Ser Leu Val Thr Met Thr Asp Ala 25 gga atg ccc gtt cca cct gga ttt gtg gtc act act gcc agc ttt gat 259 Gly Met Pro Val Pro Pro Gly Phe Val Val Thr Thr Ala Ser Phe Asp 45 gaa ttc atc cgt gaa gca ggg gtt gct gaa cac atc gat aaa ttc cta 307 Glu Phe Ile Arg Glu Ala Gly Val Ala Glu His Ile Asp Lys Phe Leu aac gat ctc gat gca gaa gat gtt aag gaa gtg gat cga gtt tct gcg 355 Asn Asp Leu Asp Ala Glu Asp Val Lys Glu Val Asp Arg Val Ser Ala 75 atc atc cgc gat gag ctg tgc agt ctt gac gtt cca gag aat gct cgt Ile Ile Arg Asp Glu Leu Cys Ser Leu Asp Val Pro Glu Asn Ala Arg ttc gca gtg cac cag gct tat cgc gat ctc atg gaa cga tgc ggt ggc 451 Phe Ala Val His Gln Ala Tyr Arg Asp Leu Met Glu Arg Cys Gly Gly 105 110 gac gtc ccg gtt gct gtc cgg tca tcg gcc act gcc gaa gat ctg ccc 499 Asp Val Pro Val Ala Val Arg Ser Ser Ala Thr Ala Glu Asp Leu Pro 547 gat gct tcc ttc gca ggg caa cag gac acc tat ctg tgg caa gtc ggt Asp Ala Ser Phe Ala Gly Gln Gln Asp Thr Tyr Leu Trp Gln Val Gly 135 140 ttg age get gte aet gaa eae ate egt aaa tge tgg get teg etg tte 595 Leu Ser Ala Val Thr Glu His Ile Arg Lys Cys Trp Ala Ser Leu Phe 150 155 act tcc cgt gcc att atc tac cgt ctg aaa aac aac atc ccc aat gag Thr Ser Arg Ala Ile Ile Tyr Arg Leu Lys Asn Asn Ile Pro Asn Glu 170 175 180 ggc ctc tcc atg gcg gta gtt gtt caa aaa atg gtc aac tct cgt gtc Gly Leu Ser Met Ala Val Val Gln Lys Met Val Asn Ser Arg Val 185 gca ggc gtg gca atc act atg aat cct tcc aac ggc gac cgc tcg aag 739 Ala Gly Val Ala Ile Thr Met Asn Pro Ser Asn Gly Asp Arg Ser Lys 200 205 atc acc atc gat tcc tca tgg ggt gtt ggt gaa atg gtg gtc tca ggt Ile Thr Ile Asp Ser Ser Trp Gly Val Gly Glu Met Val Val Ser Gly 215 220 835 gaa gtg aca cca gac aat atc ttg ctg gac aag atc acg ctg cag gtt Glu Val Thr Pro Asp Asn Ile Leu Leu Asp Lys Ile Thr Leu Gln Val

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cgc cgc agt o Arg Arg Ser 1 280									979
aag cgt gca (Lys Arg Ala (295									1027
ctg gac gct o Leu Asp Ala A 310	Asp Leu F			Asn L					1075
cgc ccg gaa a Arg Pro Glu '									1123
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Leu Asp Pro	20		25		_		30		
Thr Met Thr A	Asp Ala G	Sly Met	Pro Val 40	Pro P	ro Gly	Phe V	al Val	Thr	
Thr Ala Ser 1 50	Phe Asp G	Slu Phe 55	Ile Arg	Glu A	la Gly 60	Val A	la Glu	His	
		55		Ala G	60			•	
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Ala Glu Asp Leu Pro Asp Ala Ser Phe Ala Gly Gln Gln Asp Thr Tyr 130 135 140

Leu Trp Gln Val Gly Leu Ser Ala Val Thr Glu His Ile Arg Lys Cys 145 150 155 160

Trp Ala Ser Leu Phe Thr Ser Arg Ala Ile Ile Tyr Arg Leu Lys Asn 165 170 175

Asn Ile Pro Asn Glu Gly Leu Ser Met Ala Val Val Gln Lys Met 180 185 190

Val Asn Ser Arg Val Ala Gly Val Ala Ile Thr Met Asn Pro Ser Asn 195 200 205

Gly Asp Arg Ser Lys Ile Thr Ile Asp Ser Ser Trp Gly Val Gly Glu 210 215 220

Met Val Val Ser Gly Glu Val Thr Pro Asp Asn Ile Leu Leu Asp Lys 225 230 235 240

Ile Thr Leu Gln Val Val Ser Glu His Ile Gly Ser Lys His Ala Glu 245 250 255

Leu Ile Pro Asp Ala Thr Ser Gly Ser Leu Val Glu Lys Pro Val Asp 260 265 270

Glu Glu Arg Ala Asn Arg Arg Ser Leu Thr Asp Glu Glu Met Leu Ala 275 280 285

Val Ala Gln Met Ala Lys Arg Ala Glu Lys His Tyr Lys Cys Pro Gln 290 295 300

Asp Ile Glu Trp Ala Leu Asp Ala Asp Leu Pro Asp Gly Glu Asn Leu 305 310 315 320

Leu Leu Gln Ser Arg Pro Glu Thr Ile His Ser Asn Gly Val Lys 325 330 335

Lys Glu Thr Pro Thr Pro Gln Ala Ala Lys Thr Ile Gly Thr Phe Asp 340 345 350

Phe Ser Ser Ile Thr Val Ala Met Thr Gly Thr Lys 355 360

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Ile Ala His Leu Val His Arg Gly Ile Asp Arg Met Tyr Gly Pro Gly 50 55 60

Lys Gly Glu Asp Val Ile Tyr Tyr Ile Thr Ile Tyr Asn Glu Pro Thr 65 70 75 80

Pro Gln Pro Ala Glu Pro Glu Gly Leu Asp Val Glu Gly Leu His Lys
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Gly Ile Tyr Leu Tyr Ser Arg Gly Glu Gly Thr Gly His Glu Ala Asn 100 105 110

Ile Leu Ala Ser Gly Val Gly Met Gln Trp Ala Leu Lys Ala Ala Ser 115 120 125

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Leu Arg Asn Pro Gly Ala Asp Ala Gly Glu Ala Phe Val Thr Thr Gln
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Leu Lys Gln Thr Ser Gly Pro Tyr Val Ala Val Ser Asp Phe Ser Thr 180 185 190

Asp Leu Pro Asn Gln Ile Arg Glu Trp Val Pro Gly Asp Tyr Thr Val 195 200 205

Leu Gly Ala Asp Gly Phe Gly Phe Ser Asp Thr Arg Pro Ala Ala Arg 210 215 220

Arg Phe Phe Asn Ile Asp Ala Glu Ser Ile Val Val Ala Val Leu Asn 225 230 235 240

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cac cgt His Arg	Gly														307
atc tac Ile Tyr 70															355
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cca acc tct atg gaa cct gaa ttc cca ggc gat gag gaa atg gag aag
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Pro Thr Ser Met Glu Pro Glu Phe Pro Gly Asp Glu Glu Met Glu Lys
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cca Pro	cca Pro	ctg Leu '	gat Asp	aag Lys 325	ctt Leu	cgc Arg	tct Ser	gtc Val	cgt Arg 330	aag Lys	ggc Gly	tcc Ser	ggc Gly	aag Lys 335	cag Gln	1008
cag Gln	atc Ile	gct Ala	acc Thr	acc Thr	atg Met	gcg Ala	act Thr	gtt Val	cgt Arg	acc Thr	ttc Phe	aag Lys	gaa Glu	ctg Leu	atg Met	1056

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120

115

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- Glu His Phe Phe Gly Arg Asp Pro Arg Thr Ala Lys Leu Val Glu Asn 180 185 190
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- Asn Pro His Gly Gln Asn Tyr Val Pro Val Asp His Asp Leu Met Leu 385 390 395 400
- Ser Tyr Arg Glu Ala Pro Glu Gly Gln Ile Leu His Glu Gly Ile Asn 405 410 415
- Glu Ala Gly Ser Val Ala Ser Phe Ile Ala Ala Gly Thr Ser Tyr Ala 420 425 430
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			gcc Ala												240
			atc Ile												288
			aac Asn 100												336
Asn	Thr	Lys	atc Ile	Ile	Gln	Glu	Leu	Glu	Ser	Phe	Phe	Arg	Gly		384
			atc Ile												432
			cag Gln												480
			tac Tyr												528
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Phe Asn Arg Tyr Leu Glu Asn Arg Gly Ile Lys Asp Thr Ser Asp Gln 50 60

His Val Trp Ala Phe Leu Gly Asp Gly Glu Met Asp Glu Pro Glu Ser 65 70 75 80

Arg Gly Leu Ile Gln Gln Ala Ala Leu Asn Asn Leu Asp Asn Leu Thr 85 90 95

Phe Val Val Asn Cys Asn Leu Gln Arg Leu Asp Gly Pro Val Arg Gly
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Asn Thr Lys Ile Ile Gln Glu Leu Glu Ser Phe Phe Arg Gly Ala Gly
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Trp Ser Val Ile Lys Val Val Trp Gly Arg Glu Trp Asp Glu Leu Leu 130 135 140

Glu Lys Asp Gln Asp Gly Ala Leu Val Glu Ile Met Asn Asn Thr Ser 145 150 155 160

Asp Gly Asp Tyr Gln Thr Phe Lys Ala Asn Asp Gly Ala Tyr Val Arg 165 170 175

Glu His Phe Phe Gly Arg Asp Pro Arg Thr Ala Lys Leu Val Glu Asn 180 185 190

Met Thr Asp Glu Glu Ile Trp Lys Leu Pro Arg Gly Gly His Asp Tyr 195 200 205

Arg Lys Val Tyr Ala Ala Tyr Lys Arg Ala Leu Glu Thr Lys Asp Arg 210 215 220

Pro Thr Val Ile Leu Ala His Thr Ile Lys Gly Tyr Gly Leu Gly His 225 230 235 240

Asn Phe Glu Gly Arg Asn Ala Thr His Gln Met Lys Lys Leu Thr Leu 245 250 255

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Glu Gln Leu Glu Lys Asp Pro Tyr Leu Pro Pro Tyr Tyr His Pro Gly
275 280 285

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	gcg Ala															402
	gtc Val															450
	caa Gln															498
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	cag Gln															642
	tcc Ser															690
	gtc Val															738
	gac Asp															786
	gta Val 235															834
	gac Asp															882
	tct Ser															930
	ggc Gly															978
	gtt Val															1026
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Val Thr Val Leu Thr Tyr Ser His Trp Arg Pro Leu Gln Arg Cys Arg 320 1120 gcc gta tcg cca tgt atc acg cac tcg gtg aag gcg tgagccccat Ala Val Ser Pro Cys Ile Thr His Ser Val Lys Ala ccgtttgaag act 1133 <210> 106 <211> 341 <212> PRT <213> Corynebacterium glutamicum <400> 106 Met Ala Lys Arg Ile Val Ile Ile Gly Gly Gly Pro Ala Gly Tyr Glu Ala Ala Leu Ala Gly Ala Lys Tyr Gly Ala Glu Val Thr Val Ile Glu Asp Val Gly Val Gly Gly Ser Ala Val Thr Met Asp Cys Val Pro Ser Lys Ser Phe Ile Ala Gly Thr Gly Ile Lys Thr Asp Leu Arg Arg Ala Asp Asp Met Gly Leu Asn Arg Gly Leu Gly Lys Ala His Leu Glu Ile Asp Ala Leu Asn Ile Arg Val Lys Asp Leu Ala Lys Ala Gln Ser Glu 90 Asp Ile Leu Gly Gln Leu Gln Arg Ser Asp Val Arg Met Ile Asn Gly 100 105 Val Gly Arg Phe Asp Asp Tyr Asn Thr Lys Gln Thr Thr His Tyr Ile 120 Lys Val Thr His Ser Asp Gly Ser Glu Glu Thr Val Glu Cys Asp Leu 130 135 Val Leu Val Ala Thr Gly Ala Thr Pro Arg Ile Leu Lys Gly Ala Glu 150 Pro Asp Gly Glu Arg Ile Leu Thr Trp Arg Gln Val Tyr Asp Ile Glu Glu Leu Pro Thr His Leu Ile Val Val Gly Ser Gly Val Thr Gly Ala 180 185 Glu Phe Val Ser Ala Phe Ala Glu Leu Gly Val Lys Val Thr Met Val Ala Ser Arg Asp Arg Ile Leu Pro His Asp Asp Ala Asp Ala Asp Val Leu Glu Thr Val Leu Ala Glu Arg Gly Val Ser Leu Glu Lys His 235

Ala Arg Val Glu Ser Val Thr Arg Thr Glu Asp Gly Gly Val Cys Val Arg Thr Ala Asp Gly Arg Glu Ile Tyr Gly Ser His Ala Leu Met Thr 265 Val Gly Ser Ile Pro Asn Thr Ala Asp Leu Gly Leu Glu Asn Ile Gly 280 Val Glu Leu Ala Pro Ser Gly His Ile Lys Val Asp Arg Ser Pro Ala 290 295 Pro Thr Ser Pro Val Cys Thr Gln Gln Val Thr Val Leu Thr Tyr Ser 310 315 His Trp Arg Pro Leu Gln Arg Cys Arg Ala Val Ser Pro Cys Ile Thr 330 His Ser Val Lys Ala 340 <210> 107 <211> 1518 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (89)..(1495) <223> FRXA02853 <400> 107 aattcagcag taatcattta gacttggaac cgcttaccag tggtttcaac aatgcattca 60 cccagctcac acgtgtggag gtgccttaatg gca aag agg atc gta att atc ggc 115 Met Ala Lys Arg Ile Val Ile Ile Gly 1 ggt gga cet gea gge tat gaa gee gea ete gea gge get aaa tae ggt Gly Gly Pro Ala Gly Tyr Glu Ala Ala Leu Ala Gly Ala Lys Tyr Gly 10 15 20 gca gaa gtt acc gtt att gaa gat gtc gga gtt ggc gga tcc gca gtc Ala Glu Val Thr Val Ile Glu Asp Val Gly Val Gly Gly Ser Ala Val 30 259 acc atg gac tgt gta cct tca aag tcc ttc atc gct ggt acc ggt atc Thr Met Asp Cys Val Pro Ser Lys Ser Phe Ile Ala Gly Thr Gly Ile 45 aaa acc gac ctc cga cgt gct gat gac atg gga ctt aac cgt ggg ctt 307 Lys Thr Asp Leu Arg Arg Ala Asp Asp Met Gly Leu Asn Arg Gly Leu 60 65 gga aaa gca cac cta gaa atc gat gca ctg aac atc cgt gtg aag gac Gly Lys Ala His Leu Glu Ile Asp Ala Leu Asn Ile Arg Val Lys Asp 80 403 ctt gcg aaa gca cag tcc gaa gat atc ttg ggc cag ctg cag cgc tca Leu Ala Lys Ala Gln Ser Glu Asp Ile Leu Gly Gln Leu Gln Arg Ser

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-					gca Ala			_			_					595
_	_	_		_	att Ile 175	-	-								_	643
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_	_	_	_	_	gca Ala	_		_	_		_	-	_		-	787
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410

Val Ala Pro Thr Ala Ser Glu Leu Ile Leu Pro Ile Ala Val Ala Val

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1939

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620

600

615

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Thr Glu Thr Arg Arg Arg Thr Val Phe Asp Ala Gln Lys Trp Ile Thr 145 150 155 160

Thr His Met Arg Glu Arg His Ala Leu Gln Ser Ala Glu Pro Thr Ala 165 170 175

Arg Thr Gln Ser Lys Leu Asp Glu Ile Glu Lys Asn Ile Arg Arg Arg 180 185 190

Ile Thr Ile Leu Trp Gln Thr Ala Leu Ile Arg Val Ala Arg Pro Arg 195 200 205

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860 850 855 Leu Asp Asp Asn Pro Leu Leu Ala Arg Ser Val Gln Arg Arg Tyr Pro 870 875 Tyr Leu Leu Pro Leu Asn Val Ile Gln Val Glu Met Met Arg Arg Tyr 885 Arg Lys Gly Asp Gln Ser Glu Gln Val Ser Arg Asn Ile Gln Leu Thr 905 900 Met Asn Gly Leu Ser Thr Ala Leu Arg Asn Ser Gly 915 <210> 111 <211> 939 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(916) <223> RXN02326 <400> 111 ccaggeggac agttgtccaa cctgcgtgca caggecaccg cactgggcct tgcggatcgt 60 ttcgaactca tcgaagacaa ctacgcaagc cgttaatgag atg ctg gga cgc cca Met Leu Gly Arg Pro ace aag gte ace eca tee tee aag gtt gtt gge gae ete gea ete cae 163 Thr Lys Val Thr Pro Ser Ser Lys Val Val Gly Asp Leu Ala Leu His ctc gtt ggt gcg ggt gtg gat cca gca gac ttt gct gcc gat cca caa Leu Val Gly Ala Gly Val Asp Pro Ala Asp Phe Ala Ala Asp Pro Gln aag tac gac atc cca gac tct gtc atc gcg ttc ctg cgc ggc gag ctt 259 Lys Tyr Asp Ile Pro Asp Ser Val Ile Ala Phe Leu Arg Gly Glu Leu 45 307 ggt aac cct cca ggt ggc tgg cca gag cca ctg cgc acc cgc gca ctg Gly Asn Pro Pro Gly Gly Trp Pro Glu Pro Leu Arg Thr Arg Ala Leu gaa ggc cgc tcc gaa ggc aag gca cct ctg acg gaa gtt cct gag gaa 355 Glu Gly Arg Ser Glu Gly Lys Ala Pro Leu Thr Glu Val Pro Glu Glu gag cag gcg cac ctc gac gct gat gat tcc aag gaa cgt cgc aat agc 403 Glu Gln Ala His Leu Asp Ala Asp Asp Ser Lys Glu Arg Arg Asn Ser 90 451 ctc aac cgc ctg ctg ttc ccg aag cca acc gaa gag ttc ctc gag cac Leu Asn Arg Leu Leu Phe Pro Lys Pro Thr Glu Glu Phe Leu Glu His 105 110 115 cgt cgc cgc ttc ggc aac acc tct gcg ctg gat gat cgt gaa ttc ttc

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Asp Leu Ala Leu His Leu Val Gly Ala Gly Val Asp Pro Ala Asp Phe 20 25 30

Ala Ala Asp Pro Gln Lys Tyr Asp Ile Pro Asp Ser Val Ile Ala Phe
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Arg Thr Arg Ala Leu Glu Gly Arg Ser Glu Gly Lys Ala Pro Leu Thr
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Glu Val Pro Glu Glu Glu Gln Ala His Leu Asp Ala Asp Asp Ser Lys
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Glu Arg Arg Asn Ser Leu Asn Arg Leu Leu Phe Pro Lys Pro Thr Glu 100 105 110

Glu Phe Leu Glu His Arg Arg Phe Gly Asn Thr Ser Ala Leu Asp 115 120 125

Asp Arg Glu Phe Phe Tyr Gly Leu Val Glu Gly Arg Glu Thr Leu Ile 130 135 140

Arg Leu Pro Asp Val Arg Thr Pro Leu Leu Val Arg Leu Asp Ala Ile 145 150 155 160

Ser Glu Pro Asp Asp Lys Gly Met Arg Asn Val Val Ala Asn Val Asn 165 170 175

Gly Gln Ile Arg Pro Met Arg Val Arg Asp Arg Ser Val Glu Ser Val 180 185 190

Thr Ala Thr Ala Glu Lys Ala Asp Ser Ser Asn Lys Gly His Val Ala 195 . 200 205

Ala Pro Phe Ala Gly Val Val Thr Val Thr Val Ala Glu Gly Asp Glu 210 215 220

Val Lys Ala Gly Asp Ala Val Ala Ile Ile Glu Ala Met Lys Met Glu 225 230 235 240

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							gcc Ala 125									499
							aag Lys									547

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Tyr	Trp	Glu	Ala 265	Val	Arg	Gly	Leu	Tyr 270	Leu	Pro	Phe	Glu	Ser 275	gga Gly	Thr	931
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Ile Phe Arg Ile Phe Asp Ala Leu Asn Asp Val Ser Gln Met Arg Pro 100 , 105 110

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Met Ala Tyr Ser Gly Asp Leu Ser Asp Pro Asn Glu Lys Leu Tyr Thr 130 135 140

Leu Asp Tyr Tyr Leu Lys Met Ala Glu Glu Ile Val Lys Ser Gly Ala 145 150 155 160

His Ile Leu Ala Ile Lys Asp Met Ala Gly Leu Leu Arg Pro Ala Ala 165 170 175

Val Thr Lys Leu Val Thr Ala Leu Arg Arg Glu Phe Asp Leu Pro Val 180 185 190

His Val His Thr His Asp Thr Ala Gly Gly Gln Leu Ala Thr Tyr Phe 195 200 205

Ala Ala Gln Ala Gly Ala Asp Ala Val Asp Gly Ala Ser Gly Thr 210 215 220

Thr Val Trp His His Leu Pro Ser His Pro Leu Ser Ala Ile Val Ala 225 230 235 240

Ala Phe Ala His Thr Arg Arg Asp Thr Gly Leu Ser Leu Glu Ala Val 245 250 255

Ser Asp Leu Glu Pro Tyr Trp Glu Ala Val Arg Gly Leu Tyr Leu Pro 260 265 270

Phe Glu Ser Gly Thr Pro Gly Pro Thr Gly Arg Val Tyr Arg His Glu 275 280 285

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			_			_		-			-	-			cac His		835
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															gga Gly		931
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Ala Ile Asp Ala Val Leu Glu Thr Asn Thr Ala Val Ala Glu Val Ala 120 Met Ala Tyr Ser Gly Asp Leu Ser Asp Pro Asn Glu Lys Leu Tyr Thr 135 Leu Asp Tyr Tyr Leu Lys Met Ala Glu Glu Ile Val Lys Ser Gly Ala 150 155 His Ile Leu Ala Ile Lys Asp Met Ala Gly Leu Leu Arg Pro Ala Ala Val Thr Lys Leu Val Thr Ala Leu Arg Arg Glu Phe Asp Leu Pro Val His Val His Thr His Asp Thr Ala Gly Gly Gln Leu Ala Thr Tyr Phe 195 200 Ala Ala Ala Gln Ala Gly Ala Asp Ala Val Asp Gly Ala Ser Gly Thr 215 Thr Val Trp His His Leu Pro Ser His Pro Leu Ser Ala Ile Val Ala 230 Ala Phe Ala His Thr Arg Arg Asp Thr Gly Leu Ser Leu Glu Ala Val 250 Ser Asp Leu Glu Pro Tyr Trp Glu Ala Val Arg Gly Leu Tyr Leu Pro Phe Glu Ser Gly Thr Pro Gly Pro Thr Gly Arg Val Tyr Arg His Glu 280 Ile Pro Gly Gly Gln Leu Ser Asn Leu Arg Ala Gln Ala Thr Ala Leu 295 300 Gly Leu Ala Asp Arg Phe Glu Leu Ile Glu Asp Asn Tyr Ala Ser Arg 310

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					cgt Arg											259
					tac Tyr											307
					gct Ala 75											355
					atc Ile											403
					tac Tyr											451
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					ggt Gly 155											595
					atc Ile											643
					gtt Val											691
					gag Glu											739
					ttc Phe											787
					cat His 235											835
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480

1555

1603

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Phe Ile Ala Asp His Pro His Leu Leu Gln Ala Pro Pro Ala Asp Asp

gag cag gga cgc atc ctg gat tac ttg gca gat gtc acc gtg aac aag

475

455

470

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225 230 235 240

Leu Gly Asp His Thr Gly Glu Val Val His Leu Tyr Glu Arg Asp Cys 245 250 255

Ser Leu Gln Arg Arg His Gln Lys Val Val Glu Ile Ala Pro Ala Gln 260 265 270

His Leu Asp Pro Glu Leu Arg Asp Arg Ile Cys Ala Asp Ala Val Lys 275 280 285

Phe Cys Arg Ser Ile Gly Tyr Gln Gly Ala Gly Thr Val Glu Phe Leu 290 295 300

Val Asp Glu Lys Gly Asn His Val Phe Ile Glu Met Asn Pro Arg Ile 305 310 315 320

Gln Val Glu His Thr Val Thr Glu Glu Val Thr Glu Val Asp Leu Val 325 330 335

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Leu Thr Gln Asp Lys Ile Lys Thr His. Gly Ala Ala Leu Gln Cys Arg 355 360 365

Ile Thr Thr Glu Asp Pro Asn Asn Gly Phe Arg Pro Asp Thr Gly Thr 370 375 380

Ile Thr Ala Tyr Arg Ser Pro Gly Gly Ala Gly Val Arg Leu Asp Gly 385 390 395 400

Ala Ala Gln Leu Gly Gly Glu Ile Thr Ala His Phe Asp Ser Met Leu 405 410 415

Val Lys Met Thr Cys Arg Gly Ser Asp Phe Glu Thr Ala Val Ala Arg 420 425 430

Ala Gln Arg Ala Leu Ala Glu Phe Thr Val Ser Gly Val Ala Thr Asn 435 440 445

Ile Gly Phe Leu Arg Ala Leu Leu Arg Glu Glu Asp Phe Thr Ser Lys
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Arg Ile Ala Thr Gly Phe Ile Ala Asp His Pro His Leu Leu Gln Ala 465 470 475 480

Pro Pro Ala Asp Asp Glu Gln Gly Arg Ile Leu Asp Tyr Leu Ala Asp 485 490 495

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ggt Gly	gtg Val	cgt Arg 435	cca Pro	aag Lys	gat Asp	gtt Val	gca Ala 440	gct Ala	cct Pro	atc Ile	gat Asp	aag Lys 445	ctg Leu	cct Pro	aac Asn	1344

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Ile Asp Glu Ile Val Lys Ser Ala Glu Gly Gln Thr Tyr Pro Ile Phe 100 105 110

Val Lys Ala Val Ala Gly Gly Gly Gly Arg Gly Met Arg Phe Val Ala 115 120 125

Ser Pro Asp Glu Leu Arg Lys Leu Ala Thr Glu Ala Ser Arg Glu Ala 130 135 140

Glu Ala Ala Phe Gly Asp Gly Ala Val Tyr Val Glu Arg Ala Val Ile 145 150 155 160

Asn Pro Gln His Ile Glu Val Gln Ile Leu Gly Asp His Thr Gly Glu 165 170 175

Val Val His Leu Tyr Glu Arg Asp Cys Ser Leu Gln Arg Arg His Gln 180 185 190

Lys Val Val Glu Ile Ala Pro Ala Gln His Leu Asp Pro Glu Leu Arg . 195 200 205

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cca aag atc acc gag cta gtc gag tct gga agc ctc cag o Pro Lys Ile Thr Glu Leu Val Glu Ser Gly Ser Leu Gln I 825 830	cta aca gaa 2611 Leu Thr Glu 335
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35 40 45

Glu Pro Glu Asn Val Glu Gln Ile Arg Asp Ala Ile Ala Val Ala Val Ala Arg Gly Trp Ser Val Val Gly Arg Gly Gly Gly Ser Ser Val Ala Gly Asn Ala Ile Gly Glu Gly Leu Ile Ile Asp Thr Ser Arg Tyr Phe Asn Arg Ile Leu Asp Ile Asp Pro Val Ala Gln Thr Ala Val Val Glu 100 105 Pro Gly Val Val Cys Asp Ala Leu Arg Asp Ala Ala Ala Glu Phe Gly Leu Thr Tyr Gly Pro Asp Pro Ser Thr His Ser Arg Cys Thr Ile Gly 130 Gly Met Val Ala Asn Asn Ala Cys Gly Ser His Ser Val Ala Phe Gly Thr Ala Ala Glu Asn Leu Val Asp Val Thr Leu Met Leu Ser Asp Gly Arg Glu Val Thr Val Thr Lys Asp Gly Cys Asp Asp Ala Glu Ile Asn 185 Gln Lys Leu Thr Asp Leu Ala Ser Lys Asn Gln Asp Leu Ile Ser Lys 200 Glu Leu Gly Arg Phe Pro Arg Gln Val Ser Gly Tyr Gly Leu His Tyr Leu Ala His Asp Met Ala Lys Ala Met Ala Gly Thr Glu Gly Thr Ile 235 Gly Ile Ile Thr Arg Leu Thr Val Lys Leu Val Pro Thr Pro Lys Val Lys Ala Leu Ala Val Leu Ala Phe Asp Thr Val Phe Asp Ala Ala Arg 265 Ala Ala Ala Lys Leu Arg Leu Pro Gly Val Ala Thr Ile Glu Gly Met 275 280 Gly Gly Asp Leu Leu Ala Ala Leu Arg Ser Lys Gln Gly Gln Ser Glu Ala Gly Gln Asn Leu Pro Gly Asn Arg Ile Gly Ile Glu Ala Gly Gly 305 Trp Leu Tyr Cys Glu Thr Gly Ser Asp Thr Leu Gln Ala Ala Val Gln Ala Ala Glu Glu Val Ala Thr Ala Val Asp Thr Ile Asp Tyr Val Val 345

365

Val Ser Glu Pro Ser Glu Met Arg Glu Leu Trp Arg Ile Arg Glu Ser

360

355

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- Phe Gly His Phe Gly Glu Gly Cys Val His Val Arg Ile Ser Phe Asp 420 425 430
- Phe Ser Thr Lys Glu Gly Leu Lys Lys Phe Glu Ala Phe Met Asn Glu 435 440 445
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- Glu Met Arg Ala Leu Phe Glu Glu Phe Lys Leu Ile Phe Asp Pro Gln 485 490 495
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- Cys Val Gly Val Ser Ala Cys Arg Ser Glu Ser Asp Ala Met Cys Pro 545 550 555 560
- Ser Phe Gln Ile Thr Gly Asp Glu Val His Ser Thr Arg Gly Arg Ala 565 570 575
- Arg Leu Leu Ser Glu Met Phe Arg Gly Glu Ser Ile Ala Asp Gly Tyr 580 585 590
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- Ala Glu Phe Leu Asp Lys His Tyr Ala Gly Arg Leu Arg Pro Met Ala 625 630 635 640
- His Tyr Val Met Gly Trp Leu Pro Leu Leu Gly His Val Ala His Lys 645 650 655
- Ile Pro Leu Pro Thr Leu Ile Asp Ala Thr Met Gln Ser Ala Leu
 660 665
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Ile Ser Phe Ala His Arg Ser Leu Arg Lys Tyr Lys Pro Lys Lys Asn 690 695 700

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Asp Thr Gly Pro Ala His Ala Ala Ile Lys Thr Leu Glu Ala Leu Gly
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Tyr Asn Val Val Ile Pro Asp Gly Phe Val Cys Cys Gly Leu Thr Trp
740 745 750

His Ser Thr Gly Gln Leu Ser Met Thr Lys Lys Val Leu Glu Gln Thr 755 760 765

Ala Lys Val Met Lys Pro Tyr Leu Asp Gln Gly Leu Thr Val Val Gly 770 775 780

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Leu Gln Leu Thr Glu Ser Thr Ala Leu Thr Gln Val His Cys His Glu 835 840 845

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Ala Asp Gly Phe Ser Cys Arg Thr Gln Ile Glu Gln Gly Thr Gly Lys 915 920 925

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- Asn Val Glu Gln Ile Arg Asp Ala Ile Ala Val Ala Val Ala Arg Gly
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- Ile Gly Glu Gly Leu Ile Ile Asp Thr Ser Arg Tyr Phe Asn Arg Ile
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- Leu Asp Ile Asp Pro Val Ala Gln Thr Ala Val Val Glu Pro Gly Val
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- Gly Pro Asp Pro Ser Thr His Ser Arg Cys Thr Ile Gly Gly Met Val 130 135 140
- Ala Asn Asn Ala Cys Gly Ser His Ser Val Ala Phe Gly Thr Ala Ala 145 150 155 160
- Glu Asn Leu Val Asp Val Thr Leu Met Leu Ser Asp Gly Arg Glu Val 165 170 175
- Thr Val Thr Lys Asp Gly Cys Asp Asp Ala Glu Ile Asn Gln Lys Leu 180 185 190
- Thr Asp Leu Ala Ser Lys Asn Gln Asp Leu Ile Ser Lys Glu Leu Gly 195 200 205
- Arg Phe Pro Arg Gln Val Ser Gly Tyr Gly Leu His Tyr Leu Ala His 210 215 220
- Asp Met Ala Lys Ala Met Ala Gly Thr Glu Gly Thr Ile Gly Ile Ile 225 230 235 240
- Thr Arg Leu Thr Val Lys Leu Val Pro Thr Pro Lys Val Lys Ala Leu 245 250 255
- Ala Val Leu Ala Phe Asp Thr Val Phe Asp Ala Ala Ala Ala 260 265 270
- Lys Leu Arg Leu Pro Gly Val Ala Thr Ile Glu Gly Met Gly Gly Asp 275 280 285
- Leu Leu Ala Ala Leu Arg Ser Lys Gln Gly Gln Ser Glu Ala Gly Gln 290 295 300

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Gln

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					gaa Glu											163
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					gcc Ala											307
					atc Ile 75											355
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					ggc Gly											547
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Ala Ala Ala Phe Asp Tyr Thr Asp Gly Ala Ala Glu Ala Glu Leu Ser 50 55 60

Ile Thr Arg Ala Arg Glu Ala Phe Glu Asn Ile Glu Phe His Pro Asp 65 70 75 80

Ile Leu Lys Pro Ala Glu His Val Asp Thr Thr Thr Gln Ile Leu Gly 85 90 95

Gly Thr Ser Ser Met Pro Phe Gly Ile Ala Pro Thr Gly Phe Thr Arg
100 105 110

Leu Met Gln Thr Glu Gly Glu Ile Ala Gly Ala Gly Ala Gly Ala 115 120 125

Ala Gly Ile Pro Phe Thr Leu Ser Thr Leu Gly Thr Thr Ser Ile Glu 130 135 140

Asp Val Lys Ala Thr Asn Pro Asn Gly Arg Asn Trp Phe Gln Leu Tyr 145 150 155 160

Val Met Arg Asp Arg Glu Ile Ser Tyr Gly Leu Val Glu Arg Ala Ala 165 170 175

Lys Ala Gly Phe Asp Thr Leu Met Phe Thr Val Asp Thr Pro Ile Ala 180 185 190

Gly Tyr Arg Ile Arg Asp Ser Arg Asn Gly Phe Ser Ile Pro Pro Gln 195 200 205

Leu Thr Pro Ser Thr Val Leu Asn Ala Ile Pro Arg Pro Trp Trp 210 215 220

Ile Asp Phe Leu Thr Thr Pro Thr Leu Glu Phe Ala Ser Leu Ser Ser 225 230 235 240

Thr Gly Gly Thr Val Gly Asp Leu Leu Asn Ser Ala Met Asp Pro Thr 245 250 255

Ile Ser Tyr Glu Asp Leu Lys Val Ile Arg Glu Met Trp Pro Gly Lys 260 265 270

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														cgc Arg 180	Val	643
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														cac His		931
														gag Glu		979
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	ttc Phe															1123
	tcg Ser															1171
	ggt Gly															1219
	ccc Pro 375															1267
	cat His															1315
	aag Lys									Pro						1363
	ttc Phe															1411
	ggc Gly															1459
	atc Ile 455															1507
	aac Asn															1555
	aag Lys															1603
	gtc Val															1651
	cac His															1699
	ggt Gly															1747

535 540 545

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Pro His Lys Asp Trp Ala
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35 40 . 45

Phe Ala Val Val Arg Pro Gly Thr Leu Val Glu Met Trp Arg Ala Leu 50 55 60

Gln Val Ser Val Asp Asn Asn Leu Ile Val Ile Pro Gln Ala Ser Asn 65 70 75 80

Thr Gly Leu Thr Gly Gly Ser Gly Pro Gly Phe Gln Asp Tyr Asp Arg
85 90 95

Pro Ile Val Ile Ile Ser Thr His Arg Ile Asp Glu Val His Leu Ile 100 105 110

Asn Asp Ala Arg Glu Ala Ile Ser Leu Ala Gly Thr Pro Leu Thr His 115 120 125

Leu Thr Asp Ala Leu Ala Lys His Gln Arg Glu Pro His Ser Val Ile 130 135 140

Gly Ser Thr Ser Ile Gly Ala Ser Val Ile Gly Gly Ile Ala Asn Asn 145 150 155 160

Ser Gly Gly Ser Gln Ile Arg Lys Gly Pro Ala Phe Thr Arg Glu Ala 165 170 175

Ile Phe Ala Arg Val Asn Asp Asp Gly Lys Val Glu Leu Val Asn His
180 185 190

Leu Gly Ile Ser Leu Gly Asp Asp Pro Glu Val Ala Leu Asp Arg Leu 195 200 205 .

Gln Arg Gly Glu Trp Ser Pro Glu Asp Val Thr Pro Ala Pro Glu Asp 210 215 220

Ser Asn Glu Thr Glu Tyr Ala Glu His Leu Arg Lys Ile Val Pro Ser 225 230 235 240

Pro Ala Arg Tyr Asn Ala Asn Pro Glu Tyr Leu Phe Glu Ala Ser Gly 245 Ser Ala Gly Lys Leu Met Val Phe Ala Val Arg Thr Arg Thr Phe Pro 265 Arg Glu Val His Pro Thr Val Phe Tyr Ile Gly Thr Asn Asn Thr His 280 Glu Leu Glu Glu Ile Arg Arg Leu Phe Leu Glu Ala Asp Met Pro Leu 295 Pro Ile Ser Gly Glu Tyr Met Gly Arg Ser Ala Phe Asp Leu Ala Glu 315 Lys Tyr Gly Lys Asp Thr Phe Val Phe Leu Lys Phe Met Ser Pro Ala 330 Leu Gln Thr Arg Met Phe Ser Phe Lys Thr Trp Ala Asn Gly Leu Phe 340 Ser Lys Ile Pro Gly Ile Gly Pro Thr Phe Ala Asp Thr Val Ser Gln Ala Met Phe Ser Val Leu Pro Asn Gln Leu Pro Lys Arg Met Met Glu 375 Tyr Arg Asn Arg Phe Glu His His Leu Leu Leu Thr Val Ser Glu Ser 390 Gln Lys Ala Ala Ser Glu Lys Met Leu Lys Glu Phe Phe Ala Glu Pro Glu His Thr Gly Glu Phe Phe Ile Cys Thr Ser Asp Glu Glu Lys Ser 425 Ala Ser Leu Asn Arg Phe Gly Ala Ala Ser Ala Ala Thr Arg Tyr Ala 440 Ala Leu Lys Arg Arg His Ile Ala Gly Leu Ile Pro Ile Asp Val Ala 450 Leu Arg Arg Asp Asp Trp Asn Trp Leu Glu Val Leu Pro Glu Glu Ile 470 475 Asp Asp Gln Leu Glu Val Lys Ala Tyr Tyr Gly His Phe Phe Cys His Val Met His Gln Asp Tyr Val Ala Lys Gln Gly Val Asp Leu Glu Ala 505 Leu His Asp Arg Ile Gln His Leu Leu Glu Glu Arg Gly Ala Lys Leu 515 525 Pro Ala Glu His Asn Tyr Gly Arg Met Tyr Lys Leu Pro Glu Ser Met 535 Glu Glu His Phe Lys Glu Leu Asp Pro Thr Asn Thr Phe Asn Ala Gly 545 550

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Arg Ile Gln His Leu Leu Glu Glu His Gly Lys Lys Leu Pro Ala Glu
20 25 30

cac aac tac ggt cgc atg tac aag ctg ccg gag tcc atg gaa gag cac 144 His Asn Tyr Gly Arg Met Tyr Lys Leu Pro Glu Ser Met Glu Glu His 35 40

ttc aag gag ctc gat ccg acg aat acg ttc aac gcc ggt atc ggc ggc 192
Phe Lys Glu Leu Asp Pro Thr Asn Thr Phe Asn Ala Gly Ile Gly Gly
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His Asn Tyr Gly Arg Met Tyr Lys Leu Pro Glu Ser Met Glu Glu His 35 40 45

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Gln 65	Val	Ser			Asn 70							Gln	Ala	Ser	Asn	

70 75 80

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Pro Ile Val Ile Ile Ser Thr His Arg Ile Asp Glu Val His Leu Ile 105

Asn Asp Ala Arg Glu Ala Ile Ser Leu Ala Gly Thr Pro Leu Thr His

Leu Thr Asp Ala Leu Ala Lys His Gln Arg Glu Pro His Ser Val Ile 135

Gly Ser Thr Ser Ile Gly Ala Ser Val Ile Gly Gly Ile Ala Asn Asn 145 150 155

Ser Gly Gly Ser Gln Ile Arg Lys Gly Pro Ala Phe Thr Arg Glu Ala 170 Ile Phe Ala Arg Val Asn Asp Asp Gly Lys Val Glu Leu Val Asn His 185 Leu Gly Ile Ser Leu Gly Asp Asp Pro Glu Val Ala Leu Asp Arg Leu 200 Gln Arg Gly Glu Trp Ser Pro Glu Asp Val Thr Pro Ala Pro Glu Asp 215 Ser Asn Glu Thr Glu Tyr Ala Glu His Leu Arg Lys Ile Val Pro Ser 235 Pro Ala Arg Tyr Asn Ala Asn Pro Glu Tyr Leu Phe Glu Ala Ser Gly 245 Ser Ala Gly Lys Leu Met Val Phe Ala Val Arg Thr Arg Thr Phe Pro 265 Arg Glu Val His Pro Thr Val Phe Tyr Ile Gly Thr Asn Asn Thr His Glu Leu Glu Glu Ile Arg Arg Leu Phe Leu Glu Ala Asp Met Pro Leu Pro Ile Ser Gly Glu Tyr Met Gly Arg Ser Ala Phe Asp Leu Ala Glu 315 Lys Tyr Gly Lys Asp Thr Phe Val Phe Leu Lys Phe Met Ser Pro Ala Leu Gln Thr Arg Met Phe Ser Phe Lys Thr Trp Ala Asn Gly Leu Phe 345 Ser Lys Ile Pro Gly Ile Gly Pro Thr Phe Ala Asp Thr Val Ser Gln Ala Met Phe Ser Val Leu Pro Asn Gln Leu Pro Lys Arg Met Met Glu 375 Tyr Arg Asn Arg Phe Glu His His Leu Leu Thr Val Ser Glu Ser 385 395 390 Gln Lys Ala Ala Ser Glu Lys Met Leu Lys Glu Phe Phe Ala Glu Pro Glu His Thr Gly Glu Phe Phe Ile Cys Thr Ser Asp Glu Glu Lys Ser 420 Ala Ser Leu Asn Arg Phe Gly Ala Ala Ser Ala Ala Thr Arg Tyr Ala Ala Leu Lys Arg Arg His Ile Ala Gly Leu Ile Pro Ile Asp Val Ala 450 455 Leu Arg Arg Asp Asp Trp Asn Trp Leu Glu Val Leu Pro Glu Glu Ile 475 Asp Asp Gln Leu Glu Val Lys Ala Tyr Tyr Gly His Phe Phe Cys His

				485)				490)				495	5	
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Pr	530		a His	s Asn	ı											
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cac His	ccc Pro	acc Thr	atg Met 105	gtg Val	cgt Arg	gcc Ala	gat Asp	agt Ser 110	tgg Trp	gca Ala	cca Pro	agc Ser	act Thr 115	caa Gln	ata Ile	451
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gct ggt gga atc ggt aaa cat ctg gca gcc atg ttg aaa Ala Gly Gly Ile Gly Lys His Leu Ala Ala Met Leu Lys 135 140 145		47
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gac cat gtg gtg ttg tgc gta ccg ctt acc gca gac acc Asp His Val Val Leu Cys Val Pro Leu Thr Ala Asp Thr 185	tat cat ctg 69 Tyr His Leu 195	91
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- Gly Val Tyr Gly Gln Gln Val Ala Glu Ala Ala Met Ala Leu Leu Leu 85 90 95
- Gly Leu Ile His Met His Pro Thr Met Val Arg Ala Asp Ser Trp Ala 100 105 110
- Pro Ser Thr Gln Ile Asp Gln Gln Thr Arg Trp Leu Asp Gly Ala Thr 115 120 125
- Val Ala Ile Val Gly Ala Gly Gly Ile Gly Lys His Leu Ala Ala Met 130 135 140
- Leu Lys Pro Phe Gly Ala Lys Ser Leu Ala Val Ser Arg Thr Gly Thr 145 150 155 160
- Pro Thr Gln Asp Phe Asp Ala Thr Glu Pro Ile Ser Asn Leu His Gln
 165 170 175
- Val Leu Ala Asp Ala Asp His Val Val Leu Cys Val Pro Leu Thr Ala 180 185 190
- Asp Thr Tyr His Leu Ile Gly Lys Ala Glu Leu Lys Ala Met Gln Ser 195 200 205
- Thr Ala Ile Leu Ile Asn Val Ala Arg Gly Glu Val Val Asp Thr Glu 210 215 220
- Ala Leu Val Asp Ala Leu Asp Ala Gln Glu Ile Ser Gly Ala Gly Leu 225 230 235 240
- Asp Val Thr Asp Pro Glu Pro Leu Pro Asp Asp His Pro Leu Trp Gly 245 250 255
- Arg Ser Asn Val Ile Ile Thr Pro His Val Ala Asn Thr Leu Thr Ser 260 265 270
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Glu Glu Arg Gly Leu Asp Ile Ser Val Lys Thr Asn Ser Glu Ser Val 65 70 75 80

Thr His Arg Ser Val Leu Gln Val Lys Val Ile Thr Gly Ser Gly Ala 85 90 95

Ser Ala Thr Val Val Gly Ala Leu Thr Gly Leu Glu Arg Val Glu Lys 100 105 110

Ile Thr Arg Ile Asn Gly Arg Gly Leu Asp Leu Arg Ala Glu Gly Leu 115 120 125

Asn Leu Phe Leu Gln Tyr Thr Asp Ala Pro Gly Ala Leu Gly Thr Val 130 135 140

Gly Thr Lys Leu Gly Ala Ala Gly Ile Asn Ile Glu Ala Ala Leu 145 150 155 160

Thr Gln Ala Glu Lys Gly Asp Gly Ala Val Leu Ile Leu Arg Val Glu 165 170 175

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Asn Gly Arg Gly Leu Asp Leu Arg Ala Glu Gly Leu Asn Leu Phe Leu cag tac act gac gct cct ggt gca ctg ggt acc gtt ggt acc aag ctg Gln Tyr Thr Asp Ala Pro Gly Ala Leu Gly Thr Val Gly Thr Lys Leu ggt gct gct ggc atc aac atc gag gct gct gcg ttg act cag gct gag 192 Gly Ala Ala Gly Ile Asn Ile Glu Ala Ala Ala Leu Thr Gln Ala Glu aag ggt gac ggc gct gtc ctg atc ctg cgt gtt gag tcc gct gtc tct Lys Gly Asp Gly Ala Val Leu Ile Leu Arg Val Glu Ser Ala Val Ser 70 75 gaa gag ctg gaa gct gaa atc aac gct gag ttg ggt gct act tcc ttc Glu Glu Leu Glu Ala Glu Ile Asn Ala Glu Leu Gly Ala Thr Ser Phe 326 cag gtt gat ctt gac taattagaga tccatttgct tga Gln Val Asp Leu Asp 100 <210> 148 <211> 101 <212> PRT <213> Corynebacterium glutamicum <400> 148 Val Gly Ala Leu Thr Gly Leu Glu Arg Val Glu Lys Ile Thr Arg Ile Asn Gly Arg Gly Leu Asp Leu Arg Ala Glu Gly Leu Asn Leu Phe Leu Gln Tyr Thr Asp Ala Pro Gly Ala Leu Gly Thr Val Gly Thr Lys Leu Gly Ala Ala Gly Ile Asn Ile Glu Ala Ala Ala Leu Thr Gln Ala Glu Lys Gly Asp Gly Ala Val Leu Ile Leu Arg Val Glu Ser Ala Val Ser 65 Glu Glu Leu Glu Ala Glu Ile Asn Ala Glu Leu Gly Ala Thr Ser Phe 90 Gln Val Asp Leu Asp 100 <210> 149 <211> 604 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(604) <223> RXN03112

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gag Glu 150	gct Ala	caa Gln	gat Asp	cgt Arg	gcg Ala 155	ggt Gly	act Thr	gac Asp	att Ile	gct Ala 160	gat Asp	tct Ser	gtg Val	ctc Leu	aag Lys 165	595
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Ala	Tyr	Asp 35	Pro	Tyr	Ala	Asn	Pro 40	Ala	Arg	Ala	Ala	Gln 45	Leu	Asn	Val	
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Gly	Gly	Leu	Val 100	Asp	Glu	Gln	Ala	Leu 105	Ala	Asp	Ala	Ile	Glu 110	Ser	Gly	
His	Ile	Arg 115	Gly	Ala	Gly	Phe	Asp 120	Val	Tyr	Ser	Thr	Glu 125	Pro	Cys	Thr	
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							gtc Val									211
							aag Lys 45									259

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His Gly Gln Val Thr His Leu His Gly Arg Lys Ser Val Phe Asp Gly 65 70 75 80

Pro Thr Asp Val Leu Tyr Leu Pro Thr Gly Gln Thr Ala Thr Leu Ser 85 90 95

Gly Gln Gly Arg Val Ala Val Ala Glu Ala Pro Thr Gln Glu Pro Lys 100 105 110

Glu Trp Lys Tyr Ile Ala Pro Ala Glu Thr Pro Val Glu Leu Arg Gly
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Asn Trp Ser Ser Tyr Pro Pro His Lys His Asp Glu His Ile Pro Gly
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His Glu Ser Lys Leu Glu Glu Ile Tyr Tyr Phe Glu Ser Ala Pro Ser 180 185 190

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Tyr Ser Gly Asp Ile Ala Leu Val Pro Phe Gly Tyr His Gly Pro Ala 225 230 235 240

Val Ala Ala Pro Gly Tyr Asp Leu Tyr Tyr Leu Asn Val Met Ala Gly
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Pro Asp Pro Glu Arg Ile Trp Leu Ile Asn Asp Asp Pro Ala His Ala 260 265 270

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200 205 210

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<210> 156

<211> 288

<212> PRT

<213> Corynebacterium glutamicum

<400> 156

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20 25 30

Ile Ala Glu Leu Gly Ser Gly Glu Ser Leu Glu Leu Asn Asp Thr Gly
35 40 45

Val Glu Arg Ile Phe Ile Pro Leu Gln Gly Ser Phe Asp Val Ala His 50 55 60

His Gly Gln Val Thr His Leu His Gly Arg Lys Ser Val Phe Asp Gly 65 70 75 80

Pro Thr Asp Val Leu Tyr Leu Pro Thr Gly Gln Thr Ala Thr Leu Ser 85 90 95

Gly Gln Gly Arg Val Ala Val Ala Glu Ala Pro Thr Gln Glu Pro Lys 100 105 110

Glu Trp Lys Tyr Ile Ala Pro Ala Glu Thr Pro Val Glu Leu Arg Gly 115 120 125

Ala Gly Arg Ser Ser Arg Gln Val His Asn Phe Gly Thr Pro Glu Ala 130 135 140

Leu Asp Ala Ala Arg Leu Ile Val Cys Glu Val Ile Thr Pro Gly Glu 145 150 155 160

Asn Trp Ser Ser Tyr Pro Pro His Lys His Asp Glu His Ile Pro Gly
165 170 175

His	Glu	Ser	Lys 180	Leu	Glu	Glu	Ile	Tyr 185	Tyr	Phe	Glu	Ser	Ala 190	Pro	Ser	
Arg	Val	Gly 195	Gly	Arg	Ala	Glu	Ala 200	Ala	Glu	Gly	Ala	Phe 205	Gly	Met	Phe	
Ser	Thr 210	Tyr	Ser	Ser	Pro	Ala 215	Gly	Glu	Ile	Asp	Ile 220	Asn	Ala	Met	Val	
Tyr 225	Ser	Gly	Asp	Ile	Ala 230	Leu	Val	Pro	Phe	Gly 235	Tyr	His	Gly	Pro	Ala 240	
Val	Ala	Ala	Pro	Gly 245	Tyr	Asp	Leu	Tyr	Tyr 250	Leu	Asn	Val	Met	Ala 255	Gly	
Pro	Asp	Pro	Glu 260	Arg	Ile	Trp	Leu	Ile 265	Asn	Asp	Asp	Pro	Ala 270	His	Ala	
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Leu	Gly	Cys	Met 20	Ser	Leu	Gly	Thr	Asp 25	Tyr	Lys	Lys	Ala	Gln 30	Pro	Ile	
Ile	Glu	Ser 35	Ala	Ile	Asp	Asn	Gly 40	Ile	Thr	Tyr	Phe	Asp 45	Thr	Ala	Asp	
Ile	Tyr 50	Asp	Gln	Gly	Val	Asn 55	Glu	Glu	Ile	Val	Gly 60	Lys	Ala	Leu	Lys .	
Lys 65	Tyr	Gln	Asn	Arg	Asp 70	Asp	Ile	Val	Ile	Gly 75	Thr	Lys	Val	Gly	Asn 80	
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tca Ser	tat Tyr	ctg Leu	acc (gac Asp 10	atg Met .	gac Asp	ggc Gly	gtc Val	ctc Leu 15	atc Ile	aaa Lys	gag Glu	ggc Gly	gag Glu 20	ata Ile	163
att Ile	ccg (ggt Gly 2	gca (Ala 2	gat Asp	cgt Arg	ttt Phe	ctt (Leu (cag Gln	tct Ser	ctc Leu	acc Thr	gat Asp	aac Asn	aat Asn	gtg Val	211

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270

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265

925

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948

<210> 162

<211> 275

<212> PRT

<213> Corynebacterium glutamicum

<400> 162

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Thr Pro Arg Asp Leu Ser Ala Arg Leu Lys Thr Ser Gly Leu Asp Ile 50 55 60

Pro Pro Glu Arg Ile Trp Thr Ser Ala Thr Ala Thr Ala His Phe Leu 65 70 75 80

Lys Ser Gln Val Lys Glu Gly Thr Ala Tyr Val Val Gly Glu Ser Gly 85 90 95

Leu Thr Thr Ala Leu His Thr Ala Gly Trp Ile Leu Thr Asp Ala Asn 100 105 110

Pro Glu Phe Val Val Leu Gly Glu Thr Arg Thr Tyr Ser Phe Glu Ala 115 120 125

Ile Thr Thr Ala Ile Asn Leu Ile Leu Gly Gly Ala Arg Phe Ile Cys 130 135 140

Thr Asn Pro Asp Val Thr Gly Pro Ser Pro Ser Gly Ile Leu Pro Ala 145 150 155 160

Thr Gly Ser Val Ala Ala Leu Ile Thr Ala Ala Thr Gly Ala Glu Pro 165 170 175

Tyr Tyr Ile Gly Lys Pro Asn Pro Val Met Met Arg Ser Ala Leu Asn 180 185 190

Thr Ile Gly Ala His Ser Glu His Thr Val Met Ile Gly Asp Arg Met 195 200 205

Asp Thr Asp Val Lys Ser Gly Leu Glu Ala Gly Leu Ser Thr Val Leu 210 215 220

Val Arg Ser Gly Ile Ser Asp Asp Ala Glu Ile Arg Arg Tyr Pro Phe 225 230 235 240

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Sei	val 55	Thi	Met	Lys	Gly	Va]		. Val	. Ile	e Gly	/ Glu 65		Glu	ı Lys	a Asp	
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cct	gag Glu	gtt Val	gat Asp	ato Ile 90	: Ala	gtt Val	gac Asp	cca Pro	gtt Val 95	. Asp	ggc Gly	acc Thr	acc Thr	ctg Leu 100	atg Met	403
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gtg Val	gga Gly 135	Pro	gag Glu	gcc Ala	gca Ala	ggc Gly 140	Lys	atc Ile	gac Asp	atc Ile	gaa Glu 145	gct Ala	cca Pro	gtt Val	gcc Ala	547
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gtt Val	ctg Leu	cac His	acc Thr 265	aac Asn	gat Asp	ctg Leu	gtg Val	agc Ser 270	tcc Ser	gac Asp	aac Asn	tgc Cys	tac Tyr 275	ttc Phe	gtg Val	931
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gca Ala	aac Asn	ggc Gly	gca Ala	acc Thr	acc Thr	cgt Arg	tcc Ser	ctg Leu	gtt Val	atg Met	cgc Arg	gca Ala	aag Lys	tca Ser	ggc Gly	1027

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Arg Gln Leu Ile Asn Ser Val Thr Met Lys Gly Val Val Val Ile Gly

Glu Gly Glu Lys Asp Glu Ala Pro Met Leu Tyr Asn Gly Glu Glu Val

Gly Thr Gly Phe Gly Pro Glu Val Asp Ile Ala Val Asp Pro Val Asp

Gly Thr Thr Leu Met Ala Glu Gly Arg Pro Asn Ala Ile Ser Ile Leu 105

Ala Ala Ala Glu Arg Gly Thr Met Tyr Asp Pro Ser Ser Val Phe Tyr 120

Met Lys Lys Ile Ala Val Gly Pro Glu Ala Ala Gly Lys Ile Asp Ile 135

Glu Ala Pro Val Ala His Asn Ile Asn Ala Val Ala Lys Ser Lys Gly

Ile Asn Pro Ser Asp Val Thr Val Val Leu Asp Arg Pro Arg His 170

Ile Glu Leu Ile Ala Asp Ile Arg Arg Ala Gly Ala Lys Val Arg Leu

Ile Ser Asp Gly Asp Val Ala Gly Ala Val Ala Ala Ala Gln Asp Ser 200

Asn Ser Val Asp Ile Met Met Gly Thr Gly Gly Thr Pro Glu Gly Ile 210 215

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Let	ı Ala	a Pro	o Met	245	Asp	Phe	e Glu	u Arq	g Glr 250		s Ala	a Hi:	s As	P Ala 25	a Gly 5	
Leu	ı Val	. Leı	260	Gln	Va]	Leu	ı His	265	Asr	n Ası	p Le	ı Vai	1 Se:		r Asp	
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- Asn Ile Lys Phe Val Gln Ala Ser Met Ala Gly Ile Asp Ala Leu Val 50 60
- Lys Arg Gly Val Val Asn Glu Lys Ala Arg Trp Ala Asn Ala Gly 65 70 75 80
- Leu Tyr Ala Asp Thr Val Ala Glu Ser Thr Ile Gly Leu Ile Leu Ala 85 90 95
- Gln Met His Met His Ala Thr Thr Arg Leu Ala Lys Ser Trp Ser Val 100 105 110
- Arg Pro Glu Val Glu Asn Asn Lys Ser Trp Leu His Asp Asn Lys Thr 115 120 125
- Val Ala Ile Leu Gly Ala Gly Gly Ile Gly Val Arg Leu Leu Glu Met 130 135 140
- Leu Lys Pro Phe Asn Val Lys Thr Ile Ala Val Asn Asn Ser Gly Arg 145 150 155 160
- Pro Val Glu Gly Ala Asp Glu Thr Phe Ala Met Asp Lys Ala Glu His 165 170 175
- Val Trp Ala Glu Ala Asp Val Phe Val Leu Ile Leu Pro Leu Thr Asp 180 185 190
- Ala Thr Tyr Gln Ile Val Asn Ala Glu Thr Leu Gly Lys Met Lys Pro 195 200 205
- Ser Ala Val Val Asn Val Gly Arg Gly Pro Leu Ile Asn Thr Asp 210 215 220
- Asp Leu Val Asp Ala Leu Asn Asn Gly Thr Ile Ala Gly Ala Ala Leu 225 230 235 240
- Asp Val Thr Asp Pro Glu Pro Leu Pro Asp Ser His Pro Leu Trp Glu 245 250 255
- Met Asp Asn Val Val Ile Thr Pro His Thr Ala Asn Thr Asn Glu Arg 260 265 270
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		acc Thr														153
act Thr 35	cgt Arg	ttg Leu	gct Ala	aag Lys	tcg Ser 40	tgg Trp	agc Ser	gtg Val	cgg Arg	cct Pro 45	gag Glu	gtg Val	gaa Glu	aac Asn	aac Asn 50	201
aag Lys	tca Ser	tgg Trp	ctg Leu	cat His 55	gac Asp	aat Asn	aaa Lys	act Thr :	gtc Val 60	gct Ala	att Ile	ttg Leu	ggc Gly	gcc Ala 65	ggt Gly	249
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acc Thr	ttc Phe 100	gcc Ala	atg Met	gat Asp	aag Lys	gct Ala 105	gag Glu	cac His	gtg Val	tgg Trp	gct Ala 110	gag Glu	gct Ala	gat Asp	gtg Val	393
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gca Ala	gaa Glu	act Thr	ttg Leu	ggc Gly 135	aag Lys	atg Met	aag Lys	cct Pro	tct Ser 140	gcc Ala	gtg Val	gtg Val	gtc Val	aat Asn 145	gtg Val	489
Gly ggg	cgt Arg	ggc Gly	ccg Pro 150	ctg Leu	atc Ile	aac Asn	acc Thr	gat Asp 155	gat Asp	ctg Leu	gtg Val	gat Asp	gca Ala 160	ttg Leu	aac Asn	537
aac Asn	ggc Gly	acc Thr 165	att Ile	gcg Ala	ggt Gly	gct Ala	gcg Ala 170	ctg Leu	gac Asp	gtt Val	acc Thr	gat Asp 175	cct Pro	gag Glu	cca Pro	585

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cct cat act gca aac acg aat gag agg att cgt gct ttg acc ggc gaa Pro His Thr Ala Asn Thr Asn Glu Arg Ile Arg Ala Leu Thr Gly Glu 195 200 205 210	681
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Asn Asn Lys Ser Trp Leu His Asp Asn Lys Thr Val Ala Ile Leu Gly 50 55 60	
Ala Gly Gly Ile Gly Val Arg Leu Leu Glu Met Leu Lys Pro Phe Asn 65 70 75 80	
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Asp Val Phe Val Leu Ile Leu Pro Leu Thr Asp Ala Thr Tyr Gln Ile 115 120 125	
Val Asn Ala Glu Thr Leu Gly Lys Met Lys Pro Ser Ala Val Val 130 135 140	
Asn Val Gly Arg Gly Pro Leu Ile Asn Thr Asp Asp Leu Val Asp Ala 145 150 155 160	
Leu Asn Asn Gly Thr Ile Ala Gly Ala Ala Leu Asp Val Thr Asp Pro 165 170 175	
Glu Pro Leu Pro Asp Ser His Pro Leu Trp Glu Met Asp Asn Val Val 180 185 190	

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Met Ala Asp Gln Ala

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Asp Gly Val Ala Ser Tyr Leu Asn Asp Ser Asp Pro Glu Glu Thr Asn
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				cac His 250												883
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				gaa Glu												1075
				cca Pro 330												1123
				gtg Val												1171
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tac Tyr	ctc Leu	act Thr	ttc Phe 585	ctc Leu	gga Gly	cca Pro	cgt Arg	gtt Val 590	cct Pro	gca Ala	gag Glu	gtt Val	cac His 595	agc Ser	cag Gln	1891
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Pro Glu Ala Val Val Phe Val Asp Gly Ser Gln Ala Glu Trp Asp Arg 35 40 45

Met Ala Glu Asp Leu Val Glu Ala Gly Thr Leu Ile Lys Leu Asn Glu 50 55 60

Glu Lys Arg Pro Asn Ser Tyr Leu Ala Arg Ser Asn Pro Ser Asp Val 65 70 75 80

Ala Arg Val Glu Ser Arg Thr Phe Ile Cys Ser Glu Lys Glu Glu Asp 85 90 95

Ala Gly Pro Thr Asn Asn Trp Ala Pro Pro Gln Ala Met Lys Asp Ġlu 100 105 110

Met Ser Lys His Tyr Ala Gly Ser Met Lys Gly Arg Thr Met Tyr Val 115 120 125

Val Pro Phe Cys Met Gly Pro Ile Ser Asp Pro Asp Pro Lys Leu Gly 130 135 140

Val Gln Leu Thr Asp Ser Glu Tyr Val Val Met Ser Met Arg Ile Met 145 150 155 160

Thr Arg Met Gly Ile Glu Ala Leu Asp Lys Ile Gly Ala Asn Gly Ser 165 170 175

Phe Val Arg Cys Leu His Ser Val Gly Ala Pro Leu Glu Pro Gly Gln 180 185 190

Glu Asp Val Ala Trp Pro Cys Asn Asp Thr Lys Tyr Ile Thr Gln Phe 195 200 205

Pro Glu Thr Lys Glu Ile Trp Ser Tyr Gly Ser Gly Tyr Gly Gly Asn 210 215 220

Ala Ile Leu Ala Lys Lys Cys Tyr Ala Leu Arg Ile Ala Ser Val Met 225 230 235 240

Ala Arg Glu Glu Gly Trp Met Ala Glu His Met Leu Ile Leu Lys Leu 245 . 250 . 255

Ile Asn Pro Glu Gly Lys Ala Tyr His Ile Ala Ala Ala Phe Pro Ser 260 265 270

Ala Cys Gly Lys Thr Asn Leu Ala Met Ile Thr Pro Thr Ile Pro Gly 275 280 285

Trp Thr Ala Gln Val Val Gly Asp Asp Ile Ala Trp Leu Lys Leu Arg Glu Asp Gly Leu Tyr Ala Val Asn Pro Glu Asn Gly Phe Phe Gly Val 310 315 Ala Pro Gly Thr Asn Tyr Ala Ser Asn Pro Ile Ala Met Lys Thr Met Glu Pro Gly Asn Thr Leu Phe Thr Asn Val Ala Leu Thr Asp Asp Gly Asp Ile Trp Trp Glu Gly Met Asp Gly Asp Ala Pro Ala His Leu Ile 360 Asp Trp Met Gly Asn Asp Trp Thr Pro Glu Ser Asp Glu Asn Ala Ala 375 His Pro Asn Ser Arg Tyr Cys Val Ala Ile Asp Gln Ser Pro Ala Ala 390 395 Ala Pro Glu Phe Asn Asp Trp Glu Gly Val Lys Ile Asp Ala Ile Leu Phe Gly Gly Arg Arg Ala Asp Thr Val Pro Leu Val Thr Gln Thr Tyr Asp Trp Glu His Gly Thr Met Val Gly Ala Leu Leu Ala Ser Gly Gln 440 Thr Ala Ala Ser Ala Glu Ala Lys Val Gly Thr Leu Arg His Asp Pro 455 Met Ala Met Leu Pro Phe Ile Gly Tyr Asn Ala Gly Glu Tyr Leu Gln 470 Asn Trp Ile Asp Met Gly Asn Lys Gly Gly Asp Lys Met Pro Ser Ile 490 Phe Leu Val Asn Trp Phe Arg Arg Gly Glu Asp Gly Arg Phe Leu Trp Pro Gly Phe Gly Asp Asn Ser Arg Val Leu Lys Trp Val Ile Asp Arg 520 Ile Glu Gly His Val Gly Ala Asp Glu Thr Val Val Gly His Thr Ala 530 Lys Ala Glu Asp Leu Asp Gly Leu Asp Thr Pro Ile Glu Asp Val Lys Glu Ala Leu Thr Ala Pro Ala Glu Gln Trp Ala Asn Asp Val 565 Glu Asp Asn Ala Glu Tyr Leu Thr Phe Leu Gly Pro Arg Val Pro Ala 580 Glu Val His Ser Gln Phe Asp Ala Leu Lys Ala Arg Ile Ser Ala Ala 600

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cgt Arg	cca Pro	cgc Arg 200	Val	gag Glu	gga Gly	ttt Phe	ggt Gly 205	Leu	gaa Glu	aac Asn	act Thr	ggc Gly 210	gtt Val	aag Lys	ctc Leu	739
acc Thr	gag Glu 215	cgt Arg	ggc Gly	gca Ala	atc Ile	gag Glu 220	atc Ile	gat Asp	gat Asp	tac Tyr	atg Met 225	cgt Arg	acc Thr	aac Asn	gtc Val	787
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cca Pro	acg Thr 375	cta Leu	tct Ser	gag Glu	gca Ala	gtt Val 380	aag Lys	gaa Glu	gct Ala	Ala	cac His 385	ggt Gly	atc Ile	tct Ser	gga Gly	1267
cac His 390	atg Met	atc Ile	aac Asn	ttc Phe	taga	atco	ac c	tcgt	tggc	c ct	g		•			1305

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	_		ctt Leu					_		_		-				1027
			gct Ala													1075
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			aag Lys													1315
			ctg Leu													1363
			gcg Ala 425													1411
cgc Arg	agc Ser	cac His 440	ccc Pro	gag Glu	ctg Leu	gat Asp	gtg Val 445	gat Asp	gat Asp	gaa Glu	atc Ile	cgc Arg 450	gtg Val	cat His	ctg Leu	1459
			tat Tyr													1507
			gac Asp													1555

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240

835

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Val Asn Ala Pro Val Val Ile Asn Ala Ala Gly Pro Trp Val Ala Gln

235

Ser Gln Ala Leu Arg Gly Asn Leu Ala Glu Trp Lys Glu Thr Ile Pro Gln Asp Ala Thr Leu Val Ser Leu Ala Lys Gly Ile Glu Lys Gly Thr 105 His Leu Arg Met Ser Glu Val Ile Ala Glu Val Thr Glu Ala Asp Pro Ser Arg Ile Ala Val Leu Ser Gly Pro Asn Leu Ala Arg Glu Ile Ala 135 130 Glu Gly Gln Pro Ala Ala Thr Val Ile Ala Cys Pro Asp Glu Asn Arg Ala Lys Leu Val Gln Ala Ala Val Ala Ala Pro Tyr Phe Arg Pro Tyr Thr Asn Thr Asp Val Val Gly Thr Glu Ile Gly Gly Ala Cys Lys Asn 185 Val Ile Ala Leu Ala Cys Gly Ile Ser His Gly Tyr Gly Leu Gly Glu Asn Thr Asn Ala Ser Leu Ile Thr Arg Gly Leu Ala Glu Ile Ala Arg 215 210 Leu Gly Ala Thr Leu Gly Ala Asp Ala Lys Thr Phe Ser Gly Leu Ala 235 230 Gly Met Gly Asp Leu Val Ala Thr Cys Ser Ser Pro Leu Ser Arg Asn Arg Ser Phe Gly Glu Arg Leu Gly Gln Gly Glu Ser Leu Glu Lys Ala 265 Arg Glu Ala Thr Asn Gly Gln Val Ala Glu Gly Val Ile Ser Ser Gln Ser Ile Phe Asp Leu Ala Thr Lys Leu Gly Val Glu Met Pro Ile Thr 295 300 Gln 305 <210> 191 <211> 1809 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1786) <223> RXA01851 <400> 191 ttgtggcctt tttgcagggg aaacttattt aaataattca taagtaaaaa accgtcaatt 60 cacgatgtgg gttggcggtt ttcctattag gctcactttt atg acg agc gca cac Met Thr Ser Ala His

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Ala	Lys	Val	Phe 20		Asp	Ala	Gly	Asn 25		Val	Thr	Leu	Trp 30		Arg	
Arg	Glu	Glu 35		Ala	Ser	Thr	Ile 40		Asp	Ser	His	Glu 45		Arg	Asp	
Tyr	Leu 50		Gly	Ile	Thr	Leu 55	Pro	Glu	Ser	Leu	Gln 60	-	Thr	Ser	Ser	
Ala 65		Glu	Ala	Leu	Glu 70	Gly	Ala	Ala	Ile	Val 75		Leu	Ala	Ile	Pro 80	

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310

Ile Val Ala Leu Met Gly Arg Ser Lys Lys Ala Glu 325 330 <210> 189 <211> 1015 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1015) <223> FRXA01025 <400> 189 gggcagcagc ggcaggtttc cagraggttt ccatgcgggt ggcttggrac wtgggctaac 60 ctgaracggt taaatatcgt tttcgaaagg tgggtttcgc gtg gtt tct gta agc Val Val Ser Val Ser 163 gtg atg ggt gca ggt tcc tgg gga acc acg ttg gcc aag gtc ttc tct Val Met Gly Ala Gly Ser Trp Gly Thr Thr Leu Ala Lys Val Phe Ser 10 gat gct ggc aac gct gtg acg ttg tgg gcg agg cgg gaa gag ttg gca 211 Asp Ala Gly Asn Ala Val Thr Leu Trp Ala Arg Arg Glu Glu Leu Ala age ace ate egt gae age cat gaa aac egt gat tae ett eeg ggg att Ser Thr Ile Arg Asp Ser His Glu Asn Arg Asp Tyr Leu Pro Gly Ile 45 307 acg ttg ccg gag tcg ctg cag gtc aca tca tcg gca acg gag gct tta Thr Leu Pro Glu Ser Leu Gln Val Thr Ser Ser Ala Thr Glu Ala Leu gag ggc gca gcc att gtg gtg ttg gcg att cct tcg cag gcg ttg cgt Glu Gly Ala Ala Ile Val Val Leu Ala Ile Pro Ser Gln Ala Leu Arg 75 403 qqc aat ttq qcq qaq tqq aaa qaq acq atc ccg cag gat gcg acc ttg Gly Asn Leu Ala Glu Trp Lys Glu Thr Ile Pro Gln Asp Ala Thr Leu gtg tee ttg get aaa ggt att gaa aag gge aeg cae etg egg atg agt 451 Val Ser Leu Ala Lys Gly Ile Glu Lys Gly Thr His Leu Arg Met Ser 105 110 gaa gtg atc gcg gag gtg acg gaa gcg gat cct tca cgc atc gcg gtg 499 Glu Val Ile Ala Glu Val Thr Glu Ala Asp Pro Ser Arg Ile Ala Val . 125 120 130 547 ttq tcq qqq cca aac ctt gct cgt gag atc gcg gag ggg cag cct gca Leu Ser Gly Pro Asn Leu Ala Arg Glu Ile Ala Glu Gly Gln Pro Ala 135 140 145 gct acg gtg att gct tgc cct gat gaa aac cga gcg aaa ctt gtg cag

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Arg Glu Glu Leu Ala Ser Thr Ile Arg Asp Ser His Glu Asn Arg Asp 35 40 45

Tyr Leu Pro Gly Ile Thr Leu Pro Glu Ser Leu Gln Val Thr Ser Ser 50 55 60

Ala Thr Glu Ala Leu Glu Gly Ala Ala Ile Val Val Leu Ala Ile Pro 65 70 75 80

Ser Gln Ala Leu Arg Gly Asn Leu Ala Glu Trp Lys Glu Thr Ile Pro 85 90 95

Gln Asp Ala Thr Leu Val Ser Leu Ala Lys Gly Ile Glu Lys Gly Thr 100 105 110

His Leu Arg Met Ser Glu Val Ile Ala Glu Val Thr Glu Ala Asp Pro 115 120 125

Ser Arg Ile Ala Val Leu Ser Gly Pro Asn Leu Ala Arg Glu Ile Ala 130 135 140

Glu Gly Gln Pro Ala Ala Thr Val Ile Ala Cys Pro Asp Glu Asn Arg 145 150 155 160

Ala Lys Leu Val Gln Ala Ala Val Ala Ala Pro Tyr Phe Arg Pro Tyr 165 170 175

Thr Asn Thr Asp Val Val Gly Thr Glu Ile Gly Gly Ala Cys Lys Asn 180 185 190

Val Ile Ala Leu Ala Cys Gly Ile Ser His Gly Tyr Gly Leu Gly Glu 195 200 205

Asn Thr Asn Ala Ser Leu Ile Thr Arg Gly Leu Ala Glu Ile Ala Arg 210 215 220

Leu Gly Ala Thr Leu Gly Ala Asp Ala Lys Thr Phe Ser Gly Leu Ala 225 230 235 240

Gly Met Gly Asp Leu Val Ala Thr Cys Ser Ser Pro Leu Ser Arg Asn 245 250 255

Arg Ser Phe Gly Glu Arg Leu Gly Gln Gly Glu Ser Leu Glu Lys Ala 260 265 270

Arg Glu Ala Thr Asn Gly Gln Val Ala Glu Gly Val Ile Ser Ser Gln 275 280 285

Ser Ile Phe Asp Leu Ala Thr Lys Leu Gly Val Glu Met Pro Ile Thr 290 295 300

105 110 115

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				cgt Arg								547
				gat Asp								595
				ttc Phe								643
				gcg Ala								691
				ggc Gly 205								739
				gag Glu								787
				tct Ser								835
				ctg Leu								883
				cta Leu								931
				att Ile 285								979
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425

420

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Thr Gly Glu Pro Val Tyr Asn Ala Ile Val Trp Gln Asp Thr Arg Thr 100 105 110

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- Lys Gly Asp Leu Leu Phe Gly Thr Met Asp Thr Trp Val Leu Trp Asn 165 170 175
- Leu Thr Gly Gly Val Arg Gly Asp Asp Gly Asp Asp Ala Ile His Val 180 185 190
- Thr Asp Val Thr Asn Ala Ser Arg Thr Leu Leu Met Asp Leu Arg Thr 195 200 205
- Gln Gln Trp Asp Pro Glu Leu Cys Glu Ala Leu Asp Ile Pro Met Ser 210 215 220
- Met Leu Pro Glu Ile Arg Pro Ser Val Gly Glu Phe Arg Ser Val Arg 225 230 235 240
- His Arg Gly Thr Leu Ala Asp Val Pro Ile Thr Gly Val Leu Gly Asp 245 250 255
- Gln Gln Ala Ala Leu Phe Gly Gln Gly Gly Phe His Glu Gly Ala Ala 260 265 270
- Lys Asn Thr Tyr Gly Thr Gly Leu Phe Leu Leu Met Asn Thr Gly Thr 275 280 285
- Ser Leu Lys Ile Ser Glu His Gly Leu Leu Ser Thr Ile Ala Tyr Gln 290 295 300
- Arg Glu Gly Ser Ala Pro Val Tyr Ala Leu Glu Gly Ser Val Ser Met 305 310 315 320
- Gly Gly Ser Leu Val Gln Trp Leu Arg Asp Asn Leu Gln Leu Ile Pro 325 330 335
- Asn Ala Pro Ala Ile Glu Asn Leu Ala Arg Glu Val Glu Asp Asn Gly 340 345 350
- Gly Val His Val Val Pro Ala Phe Thr Gly Leu Phe Ala Pro Arg Trp 355 360 365
- Arg Pro Asp Ala Arg Gly Val Ile Thr Gly Leu Thr Arg Phe Ala Asn 370 375 380
- Arg Lys His Ile Ala Arg Ala Val Leu Glu Ala Asn Ala Phe Gln Thr 385 390 395 400
- Arg Glu Val Val Asp Ala Met Ala Lys Asp Ala Gly Lys Ala Leu Glu 405 410 415
- Ser Leu Arg Val Asp Gly Ala Met Val Glu Asn Asp Leu Leu Met. Gln

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gcc atg gcc aaa gac gca ggc aaa gcc ctc gaa tcc ctc cgc gtc gac Ala Met Ala Lys Asp Ala Gly Lys Ala Leu Glu Ser Leu Arg Val Asp 410 415 420	1363
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ctc ggc atc gac gtc caa cgt ctc gag gac gta gaa acc acc gcc gtc Leu Gly Ile Asp Val Gln Arg Leu Glu Asp Val Glu Thr Thr Ala Val 440 445 450	1459
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gac gag atc gaa aaa ctt att gca gtg aag aaa gtc tgg aac cct gac Asp Glu Ile Glu Lys Leu Ile Ala Val Lys Lys Val Trp Asn Pro Asp 470 475 480 485	1555
atg agc gaa gaa gag cgc gaa cgt cgc tat gcc gaa tgg aat agg gca Met Ser Glu Glu Glu Arg Glu Arg Arg Tyr Ala Glu Trp Asn Arg Ala 490 495 500	1603
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												tgg Trp		739
												cct Pro		787
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												aag Lys		979
												gga Gly		1027
I												tcc Ser		1075 /
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												cat His 355		1171
		-				_	-	_	-	_		gat Asp		1219

Val Thr Glu His Tyr Asp Val Val Leu Gly Ala Gly Pro Gly Gly

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290 295 300 Ala Ser Phe Pro Phe Ser Ala Asn Gly Lys Ala Val Gly Leu Ala Glu 310 315 Thr Asp Gly Phe Ala Lys Ile Val Ala Asp Ala Glu Phe Gly Glu Leu 330 325 Leu Gly Ala His Leu Val Gly Ala Asn Ala Ser Glu Leu Ile Asn Glu 345 Leu Val Leu Ala Gln Asn Trp Asp Leu Thr Thr Glu Glu Ile Ser Arg Ser Val His Ile His Pro Thr Leu Ser Glu Ala Val Lys Glu Ala Ala His Gly Ile Ser Gly His Met Ile Asn Phe 390 <210> 183 <211> 294 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(271) <223> RXS01261 <400> 183 gtgggtgttt ttcattttct tccactctaa aattaagtat ggaaaaccaa ccgcacccgg 60 atgcacgaca atgacccact aaacacgtat ccttgaatgc gtg act gaa cat tat Val Thr Glu His Tyr 1 gac qta qta qta ctc gga gcc ggc ccc ggt ggc tat gtc tcc gcc atc Asp Val Val Leu Gly Ala Gly Pro Gly Gly Tyr Val Ser Ala Ile cgt gca gcg cag ctt ggc aag aag gtt gct gta att gag aag cag tac Arg Ala Ala Gln Leu Gly Lys Lys Val Ala Val Ile Glu Lys Gln Tyr 30 tgg ggt ggt gtt tgc cta aac gtg ggc tgc att cct tcc aaa gtc tct 259 Trp Gly Gly Val Cys Leu Asn Val Gly Cys Ile Pro Ser Lys Val Ser gat caa aaa cgc tgaagttgcc cataccttta ccc 294 Asp Gln Lys Arg 55 <210> 184 <211> 57 <212> PRT <213> Corynebacterium glutamicum <400> 184

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- Val Thr Asp Gly Lys Asp Ala Gly Lys Thr Ile Thr Phe Asp Asp Cys 50 55 60
- Ile Ile Ala Thr Gly Ser Val Val Asn Thr Leu Arg Gly Val Asp Phe 65 70 75 80
- Ser Glu Asn Val Val Ser Phe Glu Glu Gln Ile Leu Asn Pro Val Ala 85 90 95
- Pro Lys Lys Met Val Ile Val Gly Ala Gly Ala Ile Gly Met Glu Phe 100 105 110
- Ala Tyr Val Leu Gly Asn Tyr Gly Val Asp Val Thr Val Ile Glu Phe
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- Met Asp Arg Val Leu Pro Asn Glu Asp Ala Glu Val Ser Lys Val Ile 130 135 140
- Ala Lys Ala Tyr Lys Lys Met Gly Val Lys Leu Leu Pro Gly His Ala 145 150 155 160
- Thr Thr Ala Val Arg Asp Asn Gly Asp Phe Val Glu Val Asp Tyr Gln
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- Lys Lys Gly Ser Asp Lys Thr Glu Thr Leu Thr Val Asp Arg Val Met 180 185 190
- Val Ser Val Gly Phe Arg Pro Arg Val Glu Gly Phe Gly Leu Glu Asn 195 200 205
- Thr Gly Val Lys Leu Thr Glu Arg Gly Ala Ile Glu Ile Asp Asp Tyr 210 215 220
- Met Arg Thr Asn Val Asp Gly Ile Tyr Ala Ile Gly Asp Val Thr Ala 225 230 235 240
- Lys Leu Gln Leu Ala His Val Ala Glu Ala Gln Gly Ile Val Ala Ala 245 250 255
- Glu Thr Ile Ala Gly Ala Glu Thr Gln Thr Leu Gly Asp Tyr Met Met 260 265 270
- Met Pro Arg Ala Thr Phe Cys Asn Pro Gln Val Ser Ser Phe Gly Tyr 275 280 285
- Thr Glu Glu Gln Ala Lys Glu Lys Trp Pro Asp Arg Glu Ile Lys Val

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					tca Ser											691
					att Ile											739
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ctg																840
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His	Leu	Glu	Asn 20	Thr	Met	Thr	Ala	Phe 25	Gln	Ala	Ala	Ala	Pro 30	Ala	Asp	
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Pro	Ala	Ala 115	Ala	Ala	Leu	Leu	Gln 120	Lys	Tyr	Pro	Glu	His 125	Leu	Glu	Arg	
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Tyr Tyr Asn Lys Asp Leu Trp Ala Lys Ala Gly Leu Glu Asp Arg Gly
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Pro Glu Ser Trp Glu Glu Phe Ser Glu Trp Gly Pro Lys Leu Gln Glu 20 25 30

Ala Met Asp Ser Gly Phe Ala His Gly Trp Gly Asp Ala Thr Asn Tyr 35 40 45

Leu Ser Trp Thr Phe Glu Gly Pro Met Trp Ser Leu Gly Gly Asn Tyr 50 55 60

Ser Glu Gly Trp Glu Ser Arg Leu Thr Thr Pro Glu Thr Ile Arg Ala 65 70 75 80

Val Glu Trp Leu Lys Ser Thr Val Asp Glu Gly Phe Ala Thr Val Ser 85 90 95

Thr Asp Val Thr Asn Glu Phe Ala Thr Gly Leu Ile Gly Ser Cys Ile 100 105 110

Gln Ser Thr Gly Asp Leu Ser Ser Val Ala Gly Ala Ala Ser Phe Asp 115 120 125

Trp Gly Val Ala Ala Leu Pro Asn Pro Thr Gly Glu Gly Ala Cys Pro 130 135 140

Thr Gly Gly Ala Gly Leu Gly Ile Pro Ser Gly Ile Ser Glu Gln Arg 145 150 155 160

Gln Asp Asn Ala Leu Lys Phe Ile Asp Phe Leu Thr Asn Ala Ala Asn 165 170 175

Thr Gly Tyr Trp Ser Arg Glu Thr Gly Tyr Val Pro Val Arg Lys Asp 180 185 190

Ala Ala Ser Asp Pro Asp His Ala Ala Phe Leu Glu Glu Asn Pro Ala 195 200 205

Tyr Asn Val Ala Val Glu Gln Leu Pro Asp Thr Arg Ser Gln Asp Asn 210 215 220

Phe Arg Val Leu Leu Pro Asn Gly Asp Arg Thr Ile Gly Asp Ala Leu 225 230 235 240

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tct gaa ggt tgg gag tcc cgt ctg act acc cca gag acc atc cgt gca Ser Glu Gly Trp Glu Ser Arg Leu Thr Thr Pro Glu Thr Ile Arg Ala 240

185 190 180 624 ggc tac tgg tcc cgc gag acc ggt tat gtt cca gtt cgt aag gat gct Gly Tyr Trp Ser Arg Glu Thr Gly Tyr Val Pro Val Arg Lys Asp Ala 195 200 672 gca tct gat cca gat cac gca gca ttc ctc gag gag aac cct gca tac Ala Ser Asp Pro Asp His Ala Ala Phe Leu Glu Glu Asn Pro Ala Tyr 215 210 aac gtt gca gtg gag cag ctt cct gat acc cgt tcc cag gac aac ttc 720 Asn Val Ala Val Glu Gln Leu Pro Asp Thr Arg Ser Gln Asp Asn Phe 235 230 cgc gtg ctg ctg cca aac ggt gac cgc acc atc ggt gac gca ctg gag 768 Arg Val Leu Leu Pro Asn Gly Asp Arg Thr Ile Gly Asp Ala Leu Glu 245 aag atc tgc ctg act ggt gca gac atc gat gtc acc ctg gct gag gtt Lys Ile Cys Leu Thr Gly Ala Asp Ile Asp Val Thr Leu Ala Glu Val 265 260 gag acc aag ctg aac acc atc tac acc cgc gac atc gaa cca ctt att Glu Thr Lys Leu Asn Thr Ile Tyr Thr Arg Asp Ile Glu Pro Leu Ile 275 887 taatccgagc acttcagcta cac <210> 198 <211> 288 <212> PRT <213> Corynebacterium glutamicum <400> 198 Gly Gly His Tyr Gly Leu Pro Phe Ala Arg Ser Thr Val Leu Phe Tyr Tyr Asn Lys Asp Leu Trp Ala Lys Ala Gly Leu Glu Asp Arg Gly Pro 20 Glu Ser Trp Glu Glu Phe Ser Glu Trp Gly Pro Lys Leu Gln Glu Ala Met Asp Ser Gly Phe Ala His Gly Trp Gly Asp Ala Thr Asn Tyr Leu Ser Trp Thr Phe Glu Gly Pro Met Trp Ser Leu Gly Gly Asn Tyr Ser Glu Gly Trp Glu Ser Arg Leu Thr Thr Pro Glu Thr Ile Arg Ala Val Glu Trp Leu Lys Ser Thr Val Asp Glu Gly Phe Ala Thr Val Ser Thr 105 100 Asp Val Thr Asn Glu Phe Ala Thr Gly Leu Ile Gly Ser Cys Ile Gln 120 125

Ser Thr Gly Asp Leu Ser Ser Val Ala Gly Ala Ala Ser Phe Asp Trp

135

130

Ile Leu Asp Thr Pro Thr Glu Glu Asp Phe 275 280

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Thr Glu Glu Asp Phe 280

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<211> 282

<212> PRT

<213> Corynebacterium glutamicum

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Ala Glu Leu Phe Glu Val Ser Ala Met Thr Ile His Arg Asp Leu Glu 35 40 45

Ala Leu Ala Ala Asp Asn Leu Val Glu Arg Ile Arg Gly Gly Ala Arg
50 55 60

Ser Val Ser Pro Ser Met Ser Glu Leu Ala Val Glu Gln Arg Arg His 65 70 75 80

Leu His Arg Thr Val Lys Glu Ala Leu Cys Thr Ala Ala Ala Arg Leu 85 90 95

Ile Pro Glu Gly Ala Val Val Ala Ile Asp Asp Ser Thr Thr Leu Glu 100 105 110

Ser Leu Val Glu Lys Leu Pro Gln Arg Ser Pro Ser Ala Leu Ile Thr 115 120 125

His Ser Leu Lys Thr Met Ala Asp His Arg Val Arg Ala Gly Met Ser 130 135 140

Asp Ile Arg Leu Ile Ala Cys Ala Gly Leu Tyr Phe Ala Glu Thr Asp 145 150 155 160

Ser Phe Leu Gly Lys Ala Thr Ser Ala Gln Leu Asn Glu Leu Ser Ala 165 170 175

Asp Ile Ser Phe Val Ser Thr Thr Ala Val Arg Ala Thr Gly Glu Val 180 185 190

Pro Ala Leu Phe His Pro Asp Met Glu Ala Ala Asp Thr Lys Arg Ala 195 200 205

Leu Ile Gly Ile Gly Ser Val Arg Val Leu Val Val Asp Ser Ser Lys 210 215 220

Phe Gly Ser Ala Gly Val Phe Lys Val Ala Ser Ile Glu Glu Phe Asp 225 230 235 240

His Ile Ile Ile Asp Gln Gln Cys Thr Arg Glu Gln Arg Asp Leu Leu 245 250 255

Arg Asn Ser Arg Ala Gln Ile His Val Ile Asp His Asn Gly Asp Glu 260 265 270

					att Ile											259
aat Asn	ttg Leu 55	gtg Val	gag Glu	cgc Arg	att Ile	agg Arg 60	ggt Gly	ggc Gly	gcg Ala	cgt Arg	tcg Ser 65	gtg Val	tcg Ser	ccg Pro	tcg Ser	307
					gtg Val 75											355
					act Thr											403
					gat Asp											451
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					gtg Val											547
					tat Tyr 155											595
					ttg Leu											643
					cgc Arg											691
					gct Ala											739
					gtg Val											787
					tcg Ser 235											835
					gag Glu											883
					gac Asp											931
acg	gaa	gag	gat	ttt	taag	gatgg	ict t	tggt	tctt	g ga	ıa					969

PCT/IB00/00943 WO 01/00844

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Ile Ser Glu His 115	Pro Ser Ser	Lys Gln Trp Ser 120	Ile Val Thr Asn Cy 125	7S
Leu Pro Ile Ala 130	Leu Asn Leu 135	Ala Asn Ala Gly	Leu Asp Asp Val Gl 140	Ln
Leu Leu Gly Gly 145	Ser Val Arg 150	Ala Ile Thr Gln 155	Ala Val Val Gly As 16	sp 50
Thr Ala Leu Arg	Thr Leu Ala 165	Leu Met Arg Ala 170	Asp Val Val Phe Il 175	Le
Gly Thr Asn Ala 180	Leu Thr Leu	Asp His Gly Leu 185	Ser Thr Ala Asp Se 190	er
Gln Glu Ala Ala 195	Met Lys Ser	Ala Met Ile Thr 200	Asn Ala His Lys Va 205	al
Val Val Leu Cys 210	Asp Ser Thr 215	Lys Met Gly Thr	Asp Tyr Leu Val Se 220	er
Phe Gly Ala Ile 225	Ser Asp Ile 230	Asp Val Val Val 235	Thr Asp Ala Gly Al	la 10
Pro Ala Ser Phe	Val Glu Gln 245	Leu Arg Glu Arg 250	Asp Val Glu Val Va 255	al
Ile Ala Glu				
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			atc gtt tct tat gr Ile Val Ser Tyr Va 20	
			gct gag ctt ttt g Ala Glu Leu Phe G 35	

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														cgt Arg		595
														gcg Ala 180		643
														gcc Ala		691
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														atc Ile		787
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	_	_	-	-	cgc Arg	-	-									877
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Glu	Gly	Arg	Val 20	Asn	Val	Thr	Glu	Leu 25	Ala	Gly	Arg	Phe	Asp 30	Val	Thr	
Ala	Glu	Thr 35	Ile	Arg	Arg	Asp	Leu 40	Ala	Val	Leu	Asp	Arg 45	Glu	Gly	Ile	
Val	His 50	Arg	Val	His	Gly	Gly 55	Ala	Val	Ala	Thr	Gln 60	Ser	Phe	Gln	Thr	•
Thr 65	Glu	Leu	Ser	Leu	Asp 70	Thr	Arg	Phe	Arg	Ser 75	Ala	Ser	Ser	Ala	Lys 80	
Tyr	Ser	Ile	Ala	Lys	Ala	Ala	Met	Gln	Phe	Leu	Pro	Ala	Glu	His	Gly	

Ala Ala Val Asp Leu Ala Asp Val Leu Asp Arg Arg Ile Val Leu Gly 505 Thr Leu Gly Tyr Val Gln Pro Ala Ala Val Arg Ala Thr Ala Glu Ala 520 Met Ala Gln Val Thr Gly Trp Ser Ala Glu Leu Ile Asp Ala Gln Cys 535 Gln Ser Tyr Leu Ala Lys Gln Asp Lys Ile Gln Ala Val Leu Lys Pro Tyr Arg <210> 193 <211> 900 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(877) <223> RXA01242 <400> 193 cgccggcaac caaatgaggc ttttgggcgt tggacagtga gacaatgggt aagaaattcg 60 gacatattta gtaaattggc tttttgcttt aaggagtgac atg tac gca gag gag 115 Met Tyr Ala Glu Glu cgc cgt cga cag att gcc tca tta acg gca gtt gag gga cgt gta aat Arg Arg Arg Gln Ile Ala Ser Leu Thr Ala Val Glu Gly Arg Val Asn 10 15 gtc aca gaa tta gcg ggc cga ttc gat gtc act gca gag acg att cga Val Thr Glu Leu Ala Gly Arg Phe Asp Val Thr Ala Glu Thr Ile Arg 25 cga gac ctt gcg gtg cta gac cgc gag gga att gtt cac cgc gtt cac 259 Arg Asp Leu Ala Val Leu Asp Arg Glu Gly Ile Val His Arg Val His 40 307 ggt ggc gca gta gcc acc caa tct ttc caa acc aca gag ttg agc ttg Gly Gly Ala Val Ala Thr Gln Ser Phe Gln Thr Thr Glu Leu Ser Leu 55 gat act cgt ttc agg tct gca tcg tca gca aag tac tcc att gcc aag 355 Asp Thr Arg Phe Arg Ser Ala Ser Ser Ala Lys Tyr Ser Ile Ala Lys 80 70 75 gca gcg atg cag ttc ctg ccc gct gag cat ggc gga ctg ttc ctc gat 403 Ala Ala Met Gln Phe Leu Pro Ala Glu His Gly Gly Leu Phe Leu Asp 100 90 gcg gga act act gtt act gct ttg gcc gat ctc att tct gag cat cct Ala Gly Thr Thr Val Thr Ala Leu Ala Asp Leu Ile Ser Glu His Pro

110

105

115

His Ala Glu Arg Leu Leu Ala Val Ile Lys Ala Phe Ala Ala Asp 185 Gly Gly Thr Ala Ile Asn His Ala Lys Val Thr Arg Ile Leu Arg Asn 200 Val Glu Glu Gly Arg Val Lys Gly Val Glu Val Thr Asp Gln Val Thr Asn Thr Thr His Glu Val Asn Ala Pro Val Val Ile Asn Ala Ala Gly 230 235 Pro Trp Val Ala Gln Ala Leu Gly Asp Leu Ala Glu Val Thr Lys Leu Lys Val Arg Gln Ser Lys Gly Val His Leu Leu Thr Gly Asp Leu Gly Ser Gln Ser Gly Val Phe Val Arg Gly Lys Asn Gly Lys His Val Ile Val Asn Pro Trp Met Gly Arg Thr Leu Ile Gly Pro Thr Asp Thr Met 295 Ile Asp Gly Asp Ala Asp Asp Ala Ala Ala Asp Glu Ser Asp Ile Asp Leu Leu Glu Thr Ile Asp Ser Val Arg Ala Thr Pro Leu Asp Arg 330 325 Lys Glu Ile Ile Ser Thr Leu Val Gly Val Arg Pro Leu Val Asp Asp 345 Gly Thr Asp Thr Tyr Thr Ser Ser Arg Arg Phe Asp Ile Ser Asp His Ala Asn Val Gly Ile Asp Gly Leu Val Ser Val Ser Gly Gly Lys Trp 375 Thr Thr Ser Arg Val Met Gly Tyr Lys Val Ile Glu His Val Val Glu 385 His Gln Ala Ala Val Leu Pro Pro Leu Arg His Phe Asp Ser Arg Gln 410 Met Pro Leu Ser Thr Ser Phe Gly Ala Tyr Glu Ser Val Ala Asp Ser 420 Phe Glu Ser Ala Leu Arg Ser His Pro Glu Leu Asp Val Asp Asp Glu Ile Arg Val His Leu Ala Arg Leu Tyr Gly Thr Glu His Glu Lys Val 450 455

490

Leu Asp Leu Val Ala Lys Gln Pro Asp Leu Gly Arg Arg Leu Asp Pro

Asp Asn Leu Asp Ile Ala Ala Gln Ala Val Phe Ala Val Ala Glu Glu

485

gcg gcg cag gcc gtt ttt gct gtc gcc gag gag gcg gcc gtc gac ctg Ala Ala Gln Ala Val Phe Ala Val Ala Glu Glu Ala Ala Val Asp Leu 490 495 500	1603										
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Ile Ala Arg His Ala Gln Gly Arg Gly Leu Arg Thr Val Met Phe Glu 35 40 45											
Ala Arg Asp Tyr Ser Ser Gly Thr Ser Ser Thr Thr Ser Lys Met Ile 50 55 60											
His Gly Gly Leu Arg Tyr Leu Glu Gln Tyr Asp Phe Gly Val Val Gln 65 70 75 80											
Glu Ala Val Lys Glu Arg Arg Tyr Leu Gly Ile Ala Ala Pro His Leu 85 90 95											
Val Ala Pro Arg Ser Phe Met Leu Thr Ala Phe Asp Trp Ser Glu Pro 100 105 110											
Lys Ala Pro Met Leu Gly Ala Gly Val Ala Leu Tyr Glu Thr Met Ala 115 120 125											
Trp Gln Arg Asn Gln Gly Gln Ser Lys Glu Asn His Ser Pro Arg Phe 130 135 140											
Arg Trp Ile Pro Lys Asn Ala Leu Leu Lys Glu Val Pro Trp Leu Asp 145 150 155 160											
Pro Glu Gly Leu Lys Gly Ala Trp Arg His Asp Asp Thr Leu Asn Leu 165 170 175											

Ala Gln Leu Met Met Asn Arg Gln Ile Ile Thr Gly His Leu Thr Gly 280 Ser Ala Asn Asp Thr Glu Gln Thr Met Lys Phe Ala His Leu His Gly 290 295 300 Val Lys Pro Leu Ile Glu Arg Met Pro Leu Asp Gln Ala Asm Glu Ala 310 315 Ile Ala Arg Ile Ser Ala Gly Lys Pro Arg Phe Arg Ile Val Leu Glu 325 330 Pro Asn Ser <210> 217 <211> 1641 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1618) <223> RXA02539 <400> 217 ggctgctaag cgtgcgaatg tgcgcgttgt cacaatcgtt gaccaagtgt cacctgacgc 60 acaggtagtg ctcaggtgga ggtggcccaa aggagaccca atg act gtc tac gca Met Thr Val Tyr Ala aat cca gga acc gaa ggc tcg atc gtt aac tat gaa aag cgc tac gag Asn Pro Gly Thr Glu Gly Ser Ile Val Asn Tyr Glu Lys Arg Tyr Glu 10 15 aac tac att ggt ggc aag tgg gtt cca ccg gta gag ggc cag tac ctt 211 Asn Tyr Ile Gly Gly Lys Trp Val Pro Pro Val Glu Gly Gln Tyr Leu 25 30 .gag aac att toa cot gto act ggt gaa gtt tto tgt gag gto goa cgt Glu Asn Ile Ser Pro Val Thr Gly Glu Val Phe Cys Glu Val Ala Arg 40 ggc acc gca gcg gac gtg gag ctt gca ctg gat gct gca cat gca gcc 307 Gly Thr Ala Ala Asp Val Glu Leu Ala Leu Asp Ala Ala His Ala Ala 55 get gat geg tgg ggc aag act tet gte get gaa egt get etg ate etg Ala Asp Ala Trp Gly Lys Thr Ser Val Ala Glu Arg Ala Leu Ile Leu 70 75 cac cgc att gcg gac cgc atg gaa gag cac ctg gaa gaa atc gca gtt His Arg Ile Ala Asp Arg Met Glu Glu His Leu Glu Glu Ile Ala Val 90 gca gaa acc tgg gag aac ggc aag gca gtc cgt gag act ctt gct gca 451 Ala Glu Thr Trp Glu Asn Gly Lys Ala Val Arg Glu Thr Leu Ala Ala 105 110 115

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1140

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Ala Ala Ser Gly Lys His Thr Val Phe Pro Val Thr Pro Gly His Glu

Ile Ala Gly Thr Ile Ala Glu Ile Gly Glu Asn Val Ser Arg Trp Thr

Val Gly Asp Arg Val Ala Ile Gly Trp Phe Gly Gly Asn Cys Gly Asp

Cys Ala Phe Cys Arg Ala Gly Asp Pro Val His Cys Arg Glu Arg Lys 105

Ile Pro Gly Val Ser Tyr Ala Gly Gly Trp Ala Gln Asn Ile Val Val

Pro Ala Glu Ala Leu Ala Ala Ile Pro Asp Gly Met Asp Phe Tyr Glu 135

Pro Ala Pro Met Gly Cys Ala Gly Val Thr Thr Phe Asn Ala Leu Arg 155 150

Asn Leu Lys Leu Asp Pro Gly Ala Ala Val Ala Val Phe Gly Ile Gly 170

Gly Leu Val Arg Leu Ala Ile Gln Phe Ala Ala Lys Met Gly Tyr Arg 185

Thr Ile Thr Ile Ala Arg Gly Leu Glu Arg Glu Glu Leu Ala Arg Gln

Leu Gly Ala Asn His Tyr Ile Asp Ser Asn Asp Leu His Pro Gly Gln

Ala Leu Phe Glu Leu Gly Gly Ala Asp Leu Ile Leu Ser Thr Ala Ser 230

Thr Thr Glu Pro Leu Ser Glu Leu Ser Thr Gly Leu Ser Ile Gly Gly 250

Gln Leu Thr Ile Ile Gly Val Asp Gly Gly Asp Ile Thr Val Ser Ala 265

90 95 100 gca ggt gat cet gtg cat tgc aga gag cgg aag att cet ggc gtt tet

			aga Arg					451
			aat Asn 125					499
			gac Asp					547
			aat Asn					595
			ttt Phe					643
			atg Met					691
			cta Leu 205					739
			cac His					787
			tct Ser					835
			tct Ser					883
			acc Thr					931
			cac His 285					979
			cat His					1027
			gcc Ala					1075
			att Ile					1117

Lys Tyr Leu Ser Leu Leu Lys Pro His Gly Val Met Ala Val Val Gly 70 65 Leu Pro Pro Glu Lys Gln Pro Leu Ser Phe Gly Ala Leu Ile Gly Gly Gly Lys Val Leu Thr Gly Ser Asn Ile Gly Gly Ile Pro Glu Thr Gln 105 Glu Met Leu Asp Phe Cys Ala Lys His Gly Leu Gly Ala Met Ile Glu 115 Thr Val Gly Val Asn Asp Val Asp Ala Ala Tyr Asp Arg Val Val Ala 135 Gly Asp Val Gln Phe Arg Val Val Ile Asp Thr Ala Ser Phe Ala Glu 155 150 Val Glu Ala Val <210> 215 <211> 1140 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1117) <223> RXA01758 <400> 215 cccccttatt cagagtgatg gtctaccgga gaagtaccca gaccaatagc atcgaccaac 60 gatagcgcgc tcagaagttc tttagtgaaa gcagaaccaa atg ccc aaa tac att Met Pro Lys Tyr Ile gee atg cag gta tee gaa tee ggt gea eeg tta gee geg aat ete gtg Ala Met Gln Val Ser Glu Ser Gly Ala Pro Leu Ala Ala Asn Leu Val 10 15 caa cct gct ccg ttg aaa tcg agg gaa gtc cgc gtg gaa atc gct gct 211 Gln Pro Ala Pro Leu Lys Ser Arg Glu Val Arg Val Glu Ile Ala Ala 25 259 agt ggt gtg tgc cat gca gat att ggc acg gca gca gca tcg ggg aag Ser Gly Val Cys His Ala Asp Ile Gly Thr Ala Ala Ala Ser Gly Lys 40 cac act gtt ttt cct gtt acc cct ggt cat gag att gca gga acc atc 307 His Thr Val Phe Pro Val Thr Pro Gly His Glu Ile Ala Gly Thr Ile 55 gcg gaa att ggt gaa aac gta tct cgg tgg acg gtt ggt gat cgc gtt 355 Ala Glu Ile Gly Glu Asn Val Ser Arg Trp Thr Val Gly Asp Arg Val 75 70 gca atc ggt tgg ttt ggt ggc aat tgc ggt gac tgc gct ttt tgt cgt

Ala Ile Gly Trp Phe Gly Gly Asn Cys Gly Asp Cys Ala Phe Cys Arg

gct gca gcc aag Ala Ala Ala Lys					ı Arg
aag gca gaa ctt Lys Ala Glu Leu 25	Ala Lys G				
tct gat gag gat Ser Asp Glu Asp 40		-		-	
ctc aac acc att Leu Asn Thr Ile 55					
ctc aag cca cac Leu Lys Pro His 70					
cag cca ctg ago Gln Pro Leu Ser					ı Thr
gga tcc aac att Gly Ser Asn Ile 105	Gly Gly I				
tgt gca aaa cac Cys Ala Lys His 120					
gat gtt gat gca Asp Val Asp Ala 135	Ala Tyr A				
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Arg Thr Leu Ala 35	Thr Ser A	asp Glu Asp 40	Phe Phe Thr	Glu His Ala 45	Gly
Glu Phe Asp Phe		sn Thr Ile 55	Ser Ala Ser	Ile Pro Val	Asp

<212> PRT <213> Corynebacterium glutamicum <400> 212 Val Ser Ile Ser Val Lys Ala Leu Gln Lys Ser Gly Pro Glu Ala Pro Phe Glu Val Lys Ile Ile Glu Arg Arg Asp Pro Arg Ala Asp Asp Val Val Ile Asp Ile Lys Ala Ala Gly Ile Cys His Ser Asp Ile His Thr Ile Arg Asn Glu Trp Gly Glu Ala His Phe Pro Leu Thr Val Gly His Glu Ile Ala Gly Val Val Ser Ala Val Gly Ser Asp Val Thr Lys Trp Lys Val Gly Asp Arg Val Gly Val Gly Cys Leu Val Asn Ser Cys Gly 90 Glu Cys Glu Gln Cys Val Ala Gly Phe Glu Asn Asn Cys Leu Arg Gly 100 Asn Val Gly Thr Tyr Asn Ser Asn Asp Val Asp Gly Thr Ile Thr Gln 120 Gly Gly Tyr Ala Glu Lys Val Val Val Asn Glu Arg Phe Leu Cys Ser Ile Pro Glu Glu Leu Asn Phe Asp Val Ala Ala Pro Leu Leu Cys Ala 150 Gly Ile Thr Thr Tyr Ser Pro Ile Ala Arg Trp Asn Val Lys Glu Gly Asp Lys Val Ala Val Met Gly Leu Gly Gly Thr Arg Thr His Gly Cys 185 Pro Asp Arg Cys Ser Gln Gly Cys 195 <210> 213 <211> 615 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(592) <223> RXA01572 <400> 213 etgetgtgeg caggeateae cacetaetee ceaategete getggaaegt taaagaagge 60

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303

Met Gly Val Gln Ile

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														aag Lys 20		163
														atc Ile		211
														gaa Glu		259
														ggc Gly		307
														gac Asp		355
														cag Gln 100		403
														acc Thr		451
														gct Ala		499
														gaa Glu		547
														acc Thr		595
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Gly Val Asp Tyr Leu Glu Ala Ala Pro Ile Leu Cys Ala Gly Val Thr 145 150 155 160

Val Tyr Lys Ala Leu Lys Val Ser Glu Thr Arg Pro Gly Gln Phe Met 165 170 175

Val Ile Ser Gly Val Gly Gly Leu Gly His Ile Ala Val Gln Tyr Ala 180 185 190

Ala Ala Met Gly Met Arg Val Ile Ala Val Asp Ile Ala Asp Asp Lys 195 200 205

Leu Glu Leu Ala Arg Lys His Gly Ala Glu Phe Thr Val Asn Ala Arg 210 215 220

Asn Glu Asp Ser Gly Glu Ala Val Gln Lys Tyr Thr Asn Gly Gly Ala 225 230 235 240

His Gly Val Leu Val Thr Ala Val His Glu Ala Ala Phe Gly Gln Ala 245 250 255

Leu Asp Met Ala Arg Arg Ala Gly Thr Ile Val Phe Asn Gly Leu Pro 260 265 270

Pro Gly Glu Phe Pro Ala Ser Val Phe Asn Ile Val Phe Lys Gly Leu 275 280 285

Thr Ile Arg Gly Ser Leu Val Gly Thr Arg Gln Asp Leu Ala Glu Ala 290 295 300

Leu Asp Phe Phe Ala Arg Gly Leu Ile Lys Pro Thr Val Ser Glu Cys 305 310 315 320

Ser Leu Asp Glu Val Asn Gly Val Leu Asp Arg Met Arg Asn Gly Lys 325 330 335

Ile Asp Gly Arg Val Ala Ile Arg Phe 340 345

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<223> RXA01571

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	gct Ala															835
	gca Ala															883
	gca Ala															931
	tcc Ser															979
	gtg Val 295															1027
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	ggt Gly															1123
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Pro	His	Gln 35	Ala	Leu	Val	Lys	Val 40	Leu	Thr	Ser	Gly	Ile 45	Cys	His	Thr	
Asp	Leu 50	His	Ala	Leu	Glu	Gly 55	Asp	Trp	Pro •	Val	Lys 60	Pro	Glu	Pro	Pro	
Phe 65	Val	Pro	Gly	His	Glu 70	Gly	Val	Gly	Glu	Val 75	Val	Glu	Leu	Gly	Pro 80	
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Ile Ala Val Leu Gly Ala Phe Glu His Thr Met Arg Pro Leu Gly Val 20 25 30

Thr Glu Ile Ala Glu Leu Ala Asp Leu Pro Pro Ser Thr Thr His Arg
35 40 45

Leu Val Ser Glu Leu Thr Glu Gly Gly Leu Leu Ser Lys Lys Ser Asp 50 55 60

Gly Arg Tyr Gln Leu Gly Leu Arg Ile Trp Glu Leu Ala Gln Asn Thr 65 70 75 80

Gly Arg Gln Leu Arg Asp Thr Ala Arg Pro Phe Ile Gln Glu Leu Tyr 85 90 95

Ser Leu Thr Ser Glu Thr Ala Gln Leu Val Val Arg Asp Lys Asp Glu
100 105 110

Ala Leu Leu Ile Asp Arg Ala Tyr Gly Thr Lys Lys Ile Pro Arg Ser 115 120 125

Ala Arg Val Gly Gly Arg Leu Pro Leu Asn Ser Thr Ala Val Gly Lys 130 135 140

Ile Leu Leu Ala Phe Asp Glu Pro Trp Val Lys Gln Ser Tyr Leu Lys 145 150 155 160

Leu Pro Leu Asn Ala Ser Thr Pro Lys Thr Ile Val Asn Pro Asp Val 165 170 175

Leu Ala Ala Gln Leu Lys Gln Ile His Ser Gln Gly Phe Ala Ile Thr 180 185 190

His Asp Glu Gln Arg Ile Gly Gly Ala Ser Ile Ala Val Pro Val Trp 195 200 205

His Thr Gly Lys Leu Gly Ala Ala Leu Gly Leu Val Val Pro Thr Ala 210 215 220

Gln Ala Ala Asn Leu Glu Arg Tyr Leu Pro Ile Leu Gln Ala Thr Ser 225 230 235 240

Gln Arg Ile Thr Lys Ala Thr Ala Leu Ile Pro Leu Asp Thr Leu Leu 245 250 255

Ala Ser His Lys Asn Ala Glu Arg Lys Gly Asp Thr 260 265

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<213> Corynebacterium glutamicum

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	cta Leu 135													547
	gag Glu													595
	acc Thr													643
	caa Gln													691
	ggc Gly													739
	gca Ala 215													787
	cgc Arg													835
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360 365 370

tcc aag gtg aag gtg ttt gtt att cca act aat gaa gag tta gct atc
Ser Lys Val Lys Val Phe Val Ile Pro Thr Asn Glu Glu Leu Ala Ile
375
380
385

gct agg tac gcg gtg aag ttc gct tagctctcct ggttaggatc cac 1314
Ala Arg Tyr Ala Val Lys Phe Ala
390 395

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Gly Leu Val Glu Gln Ile Gly Glu Pro Asn Gly Arg Ile Val Leu Lys 35 40 45

Ile Glu Gly Glu Lys Tyr Thr Leu Glu Thr Pro Ile Ala Asp His Ser 50 55 60

Glu Gly Leu Asn Leu Ala Phe Asp Leu Met Asp Gln His Asn Cys Gly
65 70 75 80

Pro Ser Gln Leu Glu Ile Thr Ala Val Gly His Arg Val Val His Gly 85 90 95

Gly Ile Leu Phe Ser Ala Pro Glu Leu Ile Thr Asp Glu Ile Val Glu
100 105 110

Met Ile Arg Asp Leu Ile Pro Leu Ala Pro Leu His Asn Pro Ala Asn 115 120 125

Val Asp Gly Ile Asp Val Ala Arg Lys Ile Leu Pro Asp Val Pro His 130 135 140

Val Ala Val Phe Asp Thr Gly Phe Phe His Ser Leu Pro Pro Ala Ala 145 150 155 160

Ala Leu Tyr Ala Ile Asn Lys Asp Val Ala Ala Glu His Gly Ile Arg 165 170 175

Arg Tyr Gly Phe His Gly Thr Ser His Glu Phe Val Ser Lys Arg Val 180 185 190

Val Glu Ile Leu Glu Lys Pro Thr Glu Asp Ile Asn Thr Ile Thr Phe

His Leu Gly Asn Gly Ala Ser Met Ala Ala Val Gln Gly Gly Arg Ala 210 215 220

Val Asp Thr Ser Met Gly Met Thr Pro Leu Ala Gly Leu Val Met Gly 225 230 235 240

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									gca Ala							595
									atc Ile 175							643
									cgc Arg							691
									acc Thr							739
									cgt Arg							787
									atg Met							835
									tcc Ser 255							883
									aaa Lys							931
									ctg Leu							979
									aac Asn							1027
									gca Ala							1075
									aat Asn 335							1123
									gga Gly							1171
									cga Arg							1219

170

165

175

Leu Pro Ser Trp Lys Ala Leu Asn Val Ala Ser Ile Ala Asp Leu His 185 180 Thr Lys Gly Ile Lys Val Gly Cys Trp Thr Ile Arg Asp Glu Asn Ala Phe Gly Ile Ala Gln Gln Ala Gly Val Asp Tyr Ala Thr Val Ser Asp 215 Pro Ser Arg Phe Leu Ala Pro Ser Pro Ala Gly Glu Leu His Trp <210> 205 <211> 1314 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1291) <223> RXA01436 <400> 205 gcctaaacaa accagtcaac gacctttccc gtggcgcaac agtccctgac atcgtcaaca 60 cagtagecat cacageaatt caggeaggag gacgeageta atg gea ttg gea ett Met Ala Leu Ala Leu gtt ttg aac tcc ggt tca tct tcc atc aaa ttc cag ctg gtc aac ccc 163 Val Leu Asn Ser Gly Ser Ser Ser Ile Lys Phe Gln Leu Val Asn Pro 15 10 gaa aac tot goo ato gao gag ooa tat gtt tot ggt ott gtg gag cag Glu Asn Ser Ala Ile Asp Glu Pro Tyr Val Ser Gly Leu Val Glu Gln 30 att ggt gag cca aac ggc cgc atc gta ctc aaa ata gag ggt gaa aaa 259 Ile Gly Glu Pro Asn Gly Arg Ile Val Leu Lys Ile Glu Gly Glu Lys tat acc cta gag aca ccc atc gca gat cac tcc gaa ggc cta aac ctg Tyr Thr Leu Glu Thr Pro Ile Ala Asp His Ser Glu Gly Leu Asn Leu gcg ttc gat ctc atg gac cag cac aac tgt ggt cct tcc caa ctg gaa 355 Ala Phe Asp Leu Met Asp Gln His Asn Cys Gly Pro Ser Gln Leu Glu atc acc gca gtt gga cac cgc gtg gtc cac ggc gga atc ttg ttc tcc Ile Thr Ala Val Gly His Arg Val Val His Gly Gly Ile Leu Phe Ser 90 95 gca ccg gaa ctt atc act gat gaa atc gtg gaa atg atc cgc gat ctc Ala Pro Glu Leu Ile Thr Asp Glu Ile Val Glu Met Ile Arg Asp Leu 105 110 att cca ctc gca cca ctg cac aac cct gca aac gtt gac ggc att gat

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<213> Corynebacterium glutamicum

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Gln Gly Leu Val Ser Ile Ile Thr Thr Thr Arg Asp Ala Glu Leu Ser 20 25 30

Ala Glu Leu Met Ala Asp Pro Arg Leu Ala Lys Val Thr Phe Thr Gly 35 40 45

Ser Thr Asn Val Gly Arg Ile Leu Val Arg Gln Ser Ala Asp Arg Leu 50 55 60

Leu Arg Thr Ser Met Glu Leu Gly Gly Asn Ala Ala Phe Val Ile Asp
65 70 75 80

Glu Ala Ala Asp Leu Asp Glu Ala Val Ser Gly Ala Ile Ala Ala Lys 85 90 95

Leu Arg Asn Ala Gly Gln Val Cys Ile Ala Ala Asn Arg Phe Leu Val

His Glu Ser Arg Ala Ala Glu Phe Thr Ser Lys Leu Ala Thr Ala Met 115 120 125

Gln Asn Thr Pro Ile Gly Pro Val Ile Ser Ala Arg Gln Arg Asp Arg 130 135 140

Ile Ala Ala Leu Val Asp Glu Ala Ile Thr Asp Gly Ala Arg Leu Ile 145 150 155 160

Ile Gly Gly Glu Val Pro Asp Gly Ser Gly Phe Phe Tyr Pro Ala Thr 165 170 175

Ile Leu Ala Asp Val Pro Ala Gln Ser Arg Ile Val His Glu Glu Ile 180 185 190

Phe Gly Pro Val Ala Thr Ile Ala Thr Phe Thr Asp Leu Ala Glu Gly
195 200 205

Val Ala Gln Ala Asn Ser Thr Glu Phe Gly Leu Ala Ala Tyr Gly Phe 210 215 220

Ser Asn Asn Val Lys Ala Thr Gln Tyr Met Ala Glu His Leu Glu Ala 225 230 235 240

Gly Met Val Gly Ile Asn Arg Gly Ala Ile Ser Asp Pro Ala Ala Pro 245 250 255

Phe Gly Gly Ile Gly Gln Ser Gly Phe Gly Arg Glu Gly Gly Thr Glu
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					gcc Ala											384
					Gly Ggg											432
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					acc Thr											624
					tcc Ser											672
					gca Ala 230											720
					aac Asn											768
					caa Gln											816
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Gln Asp Glu Ile Phe Gly Pro Val Pro Ser Val Val Ser Tyr Gln Asp 115 120 125

Asp Glu His Ala Ile Gln Leu Ala Asn Asp Ser Glu Phe Gly Leu Gly 130 135 140

Gly Thr Val Trp Thr Ser Asp Pro Glu Arg Gly Ala Ala Leu Ala Arg 145 150 155 160

Arg Val His Thr Gly Thr Ile Gly Ile Asn Arg Tyr Ile Pro Asp Pro 165 170 175

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Gly Pro Glu Gly Leu Ala Ser Tyr Gln Glu Thr Gln Thr Ile Tyr Leu 195 200 205

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gca gaa ctc atg gct gat cct cgc ttg gct aaa gtc acc ttc act gga 144
Ala Glu Leu Met Ala Asp Pro Arg Leu Ala Lys Val Thr Phe Thr Gly
35 40 45

tca acc aac gtg gga cgc atc ctg gtc cgc caa tcc gcg gac cga ctg 192 Ser Thr Asn Val Gly Arg Ile Leu Val Arg Gln Ser Ala Asp Arg Leu 50 55 60

ctg cgc acc tcc atg gaa ctc ggc gga aat gca gct ttt gtt atc gac 240

45 50 40 307 cag atc qqa ccq atg gcg act gcc cgg cag cgt gag cgc gtg gaa tcc Gln Ile Gly Pro Met Ala Thr Ala Arg Gln Arg Glu Arg Val Glu Ser tac att tee caa gge aaa aat get gga gee ege ate aet gte ggt gge Tyr Ile Ser Gln Gly Lys Asn Ala Gly Ala Arg Ile Thr Val Gly Gly 75 age egt eea ega gat ett gae gee gga tte tte gtt gag eea aca gtg Ser Arg Pro Arg Asp Leu Asp Ala Gly Phe Phe Val Glu Pro Thr Val ttc gcc gat gta gac aat cgc gca gcc att gcc caa gat gaa atc ttc 451 Phe Ala Asp Val Asp Asn Arg Ala Ala Ile Ala Gln Asp Glu Ile Phe 105 110 gga ccg gtg ccc tct gtt gtt tcc tac caa gac gat gaa cac gcc atc Gly Pro Val Pro Ser Val Val Ser Tyr Gln Asp Asp Glu His Ala Ile 120 caa cta gcc aac gat tcc gaa ttc ggt ctc ggc gga act gtc tgg acg Gln Leu Ala Asn Asp Ser Glu Phe Gly Leu Gly Gly Thr Val Trp Thr 135 140 age gat eee gag ege get gea ttg gee ege ega gtt eae aca gga Ser Asp Pro Glu Arg Gly Ala Ala Leu Ala Arg Arg Val His Thr Gly 150 acc att ggc atc aac cgc tat atc cct gat ccc gcc gca cca ttt gga Thr Ile Gly Ile Asn Arg Tyr Ile Pro Asp Pro Ala Ala Pro Phe Gly 170 175 ggt gtg aaa aac agt ggc ctt ggc aga gaa ctc ggc ccc gaa ggt ctt Gly Val Lys Asn Ser Gly Leu Gly Arg Glu Leu Gly Pro Glu Gly Leu 190 gct tcc tac caa gaa acc caa acc att tat ctc taatccaaac tgcacctata 744 Ala Ser Tyr Gln Glu Thr Gln Thr Ile Tyr Leu 200 205 747 tat <210> 222 <211> 208 <212> PRT <213> Corynebacterium glutamicum <400> 222 Val Glu Ala Gln Phe Thr Ser Pro Leu Leu Asn Asn Gly Gln Thr Cys Phe Leu Gly Thr Arg Ile Leu Ala Pro Lys Ser Arg Tyr. Ala Glu Val 20 25 30 Val Asp Ala Phe Thr Ala Phe Ala Gly Ser Leu Gln Val Gly Val Thr

Ser Ser Pro Asp Thr Gln Ile Gly Pro Met Ala Thr Ala Arg Gln Arg

WO 01/00844

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Asp	Ala	Val 35	Ala	Ala	Gly	Pro	Ser 40	Trp	Ala	Ala	Lys	Thr 45	Pro	Arg	Glu	
Arg	Ser 50	Val	Val	Leu	Thr	Ala 55	Ile	Phe	Glu	Ala	Leu 60	Thr	Glu	Arg	Ala	
Gln 65	Glu	Leu	Ala	Glu	Ile 70	Ile	His	Leu	Glu	Ala 75	Gly	Lys	Ser	Asp	Ala 80	
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405 410 415 Asp Glu Ala Ile Arg Ile Ala Asn Asp Thr Asn Tyr Gly Leu Gly Ala 425 Gly Val Trp Ser Arg Asp Gln Asn Thr Ile Tyr Arg Ala Gly Arg Ala 440 Ile Gln Ala Gly Arg Val Trp Val Asn Gln Tyr His Asn Tyr Pro Ala 455 His Ser Ala Phe Gly Gly Tyr Lys Glu Ser Gly Ile Gly Arg Glu Asn His Leu Met Met Leu Asn His Tyr Gln Gln Thr Lys Asn Leu Leu Val 490 Ser Tyr Asp Pro Asn Pro Thr Gly Leu Phe 500 <210> 219 <211> 430 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(430) <223> RXN03061 <400> 219 ctgccaccac tggtcattgc agaggacact ctccgtgatg gtcttcaggt gttagtcgca 60 gccctagagc gcgaaaccgc gcaccagaag gtgggctaaa gtg tct ttg acc ttc Val Ser Leu Thr Phe cca gta atc aac ccc agc gat ggc tcc acc atc acc gag cta gaa aac Pro Val Ile Asn Pro Ser Asp Gly Ser Thr Ile Thr Glu Leu Glu Asn 15 cac gat tee ace cag tgg atg tee geg ete tet gat gea get gea get 211 His Asp Ser Thr Gln Trp Met Ser Ala Leu Ser Asp Ala Val Ala Ala 259 ggt cct tca tgg gct gcg aaa act ccc cgc gaa aga tcc gtg gta ctc Gly Pro Ser Trp Ala Ala Lys Thr Pro Arg Glu Arg Ser Val Val Leu acc gca atc ttc gaa gca ctg acc gaa cgc gcc caa gaa ctt gca gag 307 Thr Ala Ile Phe Glu Ala Leu Thr Glu Arg Ala Gln Glu Leu Ala Glu atc atc cac ctg gaa gct gga aaa tcc gat gca gaa gct ctt ggt gaa 355 Ile Ile His Leu Glu Ala Gly Lys Ser Asp Ala Glu Ala Leu Gly Glu 70 gtc gct tat ggt gca gaa tac ttc cgt tgg ttt gcg gaa gaa gca gtg

95

100

Val Ala Tyr Gly Ala Glu Tyr Phe Arg Trp Phe Ala Glu Glu Ala Val

90

Arg Ala Leu Ile Leu His Arg Ile Ala Asp Arg Met Glu Glu His Leu 90 Glu Glu Ile Ala Val Ala Glu Thr Trp Glu Asn Gly Lys Ala Val Arg 100 105 Glu Thr Leu Ala Ala Asp Ile Pro Leu Ala Ile Asp His Phe Arg Tyr 120 Phe Ala Gly Ala Ile Arg Ala Gln Glu Asp Arg Ser Ser Gln Ile Asp His Asn Thr Val Ala Tyr His Phe Asn Glu Pro Ile Gly Val Val Gly 150 155 Gln Ile Ile Pro Trp Asn Phe Pro Ile Leu Met Ala Thr Trp Lys Leu Ala Pro Ala Leu Ala Ala Gly Asn Ala Ile Val Met Lys Pro Ala Glu 185 Gln Thr Pro Ala Ser Ile Leu Tyr Leu Ile Asn Ile Ile Gly Asp Leu 200 Ile Pro Glu Gly Val Leu Asn Ile Val Asn Gly Leu Gly Gly Glu Ala Gly Ala Ala Leu Ser Gly Ser Asn Arg Ile Gly Lys Ile Ala Phe Thr 230 235 Gly Ser Thr Glu Val Gly Lys Leu Ile Asn Arg Ala Ala Ser Asp Lys 250 Ile Ile Pro Val Thr Leu Glu Leu Gly Gly Lys Ser Pro Ser Ile Phe . 265 Phe Ser Asp Val Leu Ser Gln Asp Asp Ala Phe Ala Glu Lys Ala Val 280 Glu Gly Phe Ala Met Phe Ala Leu Asn Gln Gly Glu Val Cys Thr Cys 295 Pro Ser Arg Ala Leu Val His Glu Ser Ile Ala Asp Glu Phe Leu Glu Leu Gly Val Lys Arg Val Gln Asn Ile Lys Leu Gly Asn Pro Leu Asp 330 Thr Glu Thr Met Met Gly Ala Gln Ala Ser Gln Glu Gln Met Asp Lys 340 Ile Ser Ser Tyr Leu Lys Ile Gly Pro Glu Glu Gly Ala Gln Thr Leu Thr Gly Gly Lys Val Asn Lys Val Asp Gly Met Glu Asn Gly Tyr Tyr 375 Ile Glu Pro Thr Val Phe Arg Gly Thr Asn Asp Met Arg Ile Phe Arg Glu Glu Ile Phe Gly Pro Val Leu Ser Val Ala Thr Phe Ser Asp Phe.

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cgt g Arg A																547
tac c Tyr H 150											Gln					595
aac t Asn P																643
gca g Ala G																691
att t Ile L																739
ctc a Leu A 2																787
ggc t Gly S 230																835
ggc a Gly L																883
ctg g Leu G																931
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		gtg Val									1459
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		atc Ile 475									1555
		acg Thr									1603
		cac His									1651
		cgt Arg									1699
		cag Gln									1747
		tac Tyr 555									1795
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		ttc Phe									1891
		ggt Gly									1939
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					cac His 75											355
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Leu					cag Gln											547
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547

125

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120

135

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- Tyr Glu Glu Val Ile Ala Arg Phe Ser Lys Ala Ala Lys Ala Met Ser 305 310 315 320
- Ile Gly Ala Gly Phe Glu Trp Lys Tyr Glu Met Gly Ser Leu Ile Asn 325 330 335
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- Ile Arg Asn Ile Ala Glu Gln Arg Trp Met Ser Met Arg Gly Pro Ala

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Met Ile Lys Arg Leu Pro S Thr Leu Leu Asp Leu The 20 The 35 The 3	ar Ala Asn Ala 25 ar Gly Glu Thr 40 au His Ala Phe 55 ar Thr Ala Val 0 al Leu Lys Asn by Lys Asn Arg 105	Gln Asp Ala Ala Leu Gly Phe Gly 45 Ala Leu Ser Arg 60 Glu Arg Lys Lys 75 Arg Glu Leu Leu 90 Ala Ser Ala Ala	Lys Val Glu 30 Phe Asp Gly Ala Ala Gln Ile Phe Leu 80 Met Asp Ile 95 Asp Glu Val 110	
Met Ile Lys Arg Leu Pro S Thr Leu Leu Asp Leu Th 20 Val Ile Ala Pro Phe Th 35 Asp Glu Gln Asp Val Gl 57 Lys Lys Trp Val His Th 7 Lys Val His Asp Leu Va 85 Val Gln Leu Glu Thr Gl 100 Leu Asp Val Ala Ile Th	r Ala Asn Ala 25 r Gly Glu Thr 40 u His Ala Phe 55 r Thr Ala Val 0 l Leu Lys Asn y Lys Asn Arg 105 r Thr Arg Phe 120	Gln Asp Ala Ala Leu Gly Phe Gly 45 Ala Leu Ser Arg 60 Glu Arg Lys Lys 75 Arg Glu Leu Leu 90 Ala Ser Ala Ala Tyr Ala Asn Asn 125	Lys Val Glu 30 Phe Asp Gly Ala Ala Gln Ile Phe Leu 80 Met Asp Ile 95 Asp Glu Val 110 Ala Gly Lys	

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					gat Asp											787
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499

547

595

639

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Gln 65	Ile	Thr	Lys	Gln	Leu 70	Asn	Lys	Leu	Ile	Pro 75	Val	Leu	Lys	Val	Val 80
Arg	Leu	Asp	Glu	Glu 85	Thr	Thr	Ile	Ala	Arg 90	Ala	Ile	Met	Leu	Val 95	Lys
Val	Ser	Ala	Asp 100	Ser	Thr	Asn	Arg	Pro 105	Gln	Ile	Val	Asp	Ala 110	Ala	Asn
Ile	Phe	Arg 115	Ala	Arg	Val	Val	Asp 120	Val	Ala	Pro	Asp	Ser 125	Val	Val	Ile
Glu	Ser 130	Thr	Gly	Thr	Pro	Gly 135	Lys	Leu	Arg	Ala	Leu 140	Leu	Asp	Val	Met
Glu 145	Pro	Phe	Gly	Ile	Arg 150	Glu	Leu	Ile	Gln	Ser 155	Gly	Gln	Ile	Ala	Leu 160
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gaa Gli	a tto 1 Leu	gat Asp	ttc Phe	yal Val	. Trp	cca Pro	cca Pro	aag Lys	ato Ile 175	Asp	ctg Leu	cca Pro	ggc	tac Tyr 180	cgc Arg	643
cca Pro	gtt Val	tca Ser	aca Thr 185	Pro	cat His	gct Ala	cgc Arg	cag Gln 190	Ile	gag Glu	cag Gln	gca Ala	gtc Val 195	Lys	ctg Leu	691
ato Ile	ggt Gly	gag Glu 200	Ala	aag Lys	aag Lys	ccc Pro	gtc Val 205	Leu	tac Tyr	gtt Val	ggt Gly	ggt Gly 210	ggc Gly	gta Val	atc	739
aag Lys	gct Ala 215	gac Asp	gca Ala	cac His	gaa Glu	gag Glu 220	ctt Leu	cgt Arg	gcg Ala	ttc Phe	gct Ala 225	gag Glu	tac Tyr	acc Thr	ggc Gly	787
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Glu Ser Thr Glu Ala 370

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Met Thr Gly Ala Lys

1 5

gca att gtt cga tcg ctc gag gag ctt aac gcc gac atc gtg ttc ggt 163 Ala Ile Val Arg Ser Leu Glu Glu Leu Asn Ala Asp Ile Val Phe Gly 10 15 20

att cct ggt ggt gcg gtg cta ccg gtg tat gac ccg ctc tat tcc tcc 211
Ile Pro Gly Gly Ala Val Leu Pro Val Tyr Asp Pro Leu Tyr Ser Ser
25 30 35

aca aag gtg cgc cac gtc ttg gtg cgc cac gag cag ggc gca ggc cac 259
Thr Lys Val Arg His Val Leu Val Arg His Glu Gln Gly Ala Gly His
40 45 50

gca gca acc ggc tac gcg cag gtt act gga cgc gtt ggc gtc tgc att 307 Ala Ala Thr Gly Tyr Ala Gln Val Thr Gly Arg Val Gly Val Cys Ile 55 . 60 65

gca acc tct ggc cca gga gca acc aac ttg gtt acc cca atc gct gat 355
Ala Thr Ser Gly Pro Gly Ala Thr Asn Leu Val Thr Pro Ile Ala Asp
70 75 80 85

gca aac ttg gac tcc gtt ccc atg gtt gcc atc acc ggc cag gtc gga 403 Ala Asn Leu Asp Ser Val Pro Met Val Ala Ile Thr Gly Gln Val Gly

agt ggc ctg ctg ggt acc gac gct ttc cag gaa gcc gat atc cgc ggc 451 Ser Gly Leu Leu Gly Thr Asp Ala Phe Gln Glu Ala Asp Ile Arg Gly 105 110 115

atc acc atg cca gtg acc aag cac, aac ttc atg gtc acc aac cct aac 499

Ile Thr Met Pro Val Thr Lys His Asn Phe Met Val Thr Asn Pro Asn

120 125 130

gac att cca cag gca ttg gct gag gca ttc cac ctc gcg att act ggt 54° Asp Ile Pro Gln Ala Leu Ala Glu Ala Phe His Leu Ala Ile Thr Gly 135 140 145

His Glu Leu His Met Gly Met Pro Gly Met His Gly Thr Val Ser Ala 20 25 30

- Val Gly Ala Leu Gln Arg Ser Asp Leu Leu Ile Ala Ile Gly Ser Arg
 35 40 45
- Phe Asp Asp Arg Val Thr Gly Asp Val Asp Thr Phe Ala Pro Asp Ala 50 55 60
- Lys Ile Ile His Ala Asp Ile Asp Pro Ala Glu Ile Gly Lys Ile Lys
 65 70 75 80
- Gln Val Glu Val Pro Ile Val Gly Asp Ala Arg Glu Val Leu Ala Arg 85 90 95
- Leu Leu Glu Thr Thr Lys Ala Ser Lys Ala Glu Thr Glu Asp Ile Ser 100 105 110
- Glu Trp Val Asp Tyr Leu Lys Gly Leu Lys Ala Arg Phe Pro Arg Gly 115 120 125
- Tyr Asp Glu Gln Pro Gly Asp Leu Leu Ala Pro Gln Phe Val Ile Glu 130 135 140
- Thr Leu Ser Lys Glu Val Gly Pro Asp Ala Ile Tyr Cys Ala Gly Val 145 150 155 160
- Gly Gln His Gln Met Trp Ala Ala Gln Phe Val Asp Phe Glu Lys Pro 165 170 175
- Arg Thr Trp Leu Asn Ser Gly Gly Leu Gly Thr Met Gly Tyr Ala Val
- Pro Ala Ala Leu Gly Ala Lys Ala Gly Ala Pro Asp Lys Glu Val Trp 195 200 205
- Ala Ile Asp Gly Asp Gly Cys Phe Gln Met Thr Asn Gln Glu Leu Thr 210 220
- Thr Ala Ala Val Glu Gly Phe Pro Ile Lys Ile Ala Leu Ile Asn Asn 225 230 235 240
- Gly Lys Pro Gly Ala Trp Val Arg Gln Trp Gln Thr Leu Phe Tyr Glu 245 250 255
- Gly Arg Tyr Ser Asn Thr Lys Leu Arg Asn Gln Gly Glu Tyr Met Pro 260 265 270
- Asp Phe Val Thr Leu Ser Glu Gly Leu Gly Cys Val Ala Ile Arg Val 275 280 285
- Thr Lys Ala Glu Glu Val Leu Pro Ala Ile Gln Lys Ala Arg Glu Ile 290 295 300
- Asn Asp Arg Pro Val Val Ile Asp Phe Ile Val Gly Glu Asp Ala Gln 305 310 310 320
- Val Trp Pro Met Val Ser Ala Gly Ser Ser Asn Ser Asp Ile Gln Tyr 325 330 335
- Ala Leu Gly Leu Arg Pro Phe Phe Asp Gly Asp Glu Ser Ala Ala Glu

Arg Thr Trp	Leu Asn 180	Ser Gly	/ Gly	Leu 185	Gly	Thr	Met	Gly	Tyr 190	Ala	Val	
cct gcg gcc Pro Ala Ala 195	Leu Gly											624
gct atc gac Ala Ile Asp 210			Phe									672
acc gcc gca Thr Ala Ala 225												720
gga aaa cct Gly Lys Pro		Trp Val										768
gga cgg tac Gly Arg Tyr				_		_				_		816
gac ttt gtt Asp Phe Val 275						-		_		_		864
acc aaa gcg Thr Lys Ala 290			Pro									912
aac gac cgc Asn Asp Arg 305												960
gta tgg cca Val Trp Pro												1008
gca ctc gga Ala Leu Gly												1056
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Tyr Leu Lys Asn Gln Ala Leu Gln Arg Pro Leu Leu Gly

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576

cgc acc tgg ctc aac tcc ggt gga ctg ggc acc atg ggc tac gca gtt

295 300 290 Ile Gly Ile Gly Thr Arg Tyr Ser Asp Phe Thr Thr Ala Ser Arg Thr 310 315 Ala Phe Gln Asn Pro Asp Val Thr Phe Ile Asn Ile Asn Val Ala Ser 330 Phe Asp Ala Tyr Lys His Gly Thr Gln Leu Pro Val Ile Ala Asp Ala 345 Arg Glu Ala Ile Val Glu Leu Ala Glu Ala Leu Gln Gly Phe Thr Val Ala Glu Asp Tyr Ala Gln Arg Ile Ala Lys Glu Lys Ala Ala Trp Asp 375 Ala Glu Val Asp Lys Ser Phe Ala Pro Ser Gly Leu Ala Leu Pro Gly 385 Gln Pro Glu Ile Ile Gly Ala Val Gln Ala Ser Thr Ser Glu Lys Asp 410 Val Ile Val Gln Ala Ala Gly Ser Leu Pro Gly Asp Leu His Lys Leu Trp Arg Val Arg Asp Ala Leu Gly Tyr His Val Glu Tyr Ala Phe Ser Cys Met Gly Tyr Glu Ile Ala Gly Gly Ile Gly Ala Lys Arg Gly Leu Asp Ala Ala Gly Asp Asp Arg Asp Val Val Ile Met Val Gly Asp Gly Ser Tyr Leu Met Leu Asn Thr Glu Leu Val Thr Ala Val Ala Glu Gly 490 Ile Lys Val Ile Val Val Leu Ile Gln Asn His Gly Tyr Ala Ser Ile 505 500 Gly His Leu Ser Glu Thr Val Gly Ser Gln Arg Phe Gly Thr Trp Tyr Arg Glu Tyr Asp Ala Glu Ala Lys Asn Phe Gln Gly Glu Gln Ile Leu 530 Pro Val Asp Leu Ala Met Asn Ala Arg Ser Tyr Gly Met Asp Val Ile Glu Val Glu Pro Ser Ala Asn Ala Ile Glu Asp Leu Lys Ala Ala Met 565 Ala Thr Ala Lys Ala Ser Glu Lys Ser Thr Phe Ile His Ile Asn Ser 585 Asp Pro Leu Ile Tyr Ala Pro Asp Gly Ala Gly Trp Trp Asp Val Pro 605

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615

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- Pro Gly Met Phe Gly Ile Phe Gly His Gly Asn Val Ala Gly Ile Gly 35 40 45
- Gln Ala Leu Lys Gln Tyr Asn Val Glu Gln Pro Glu Leu Met Pro Tyr
 50 60
- Tyr Gln Ala Arg Asn Glu Gln Ala Met Val His Gln Ser Val Gly Tyr 65 70 75 80
- Ala Arg Met His Arg Arg Gly Thr Tyr Ala Ser Ala Ala Ser Val 85 90 95
- Gly Pro Gly Ala Thr Asn Leu Leu Thr Gly Ala Ala Leu Ala Thr Thr 100 105 110
- Asn Arg Leu Pro Ala Leu Leu Pro Ser Asp Thr Phe Ala Thr Arg 115 120 125
- Val Ala Asp Pro Val Leu Gln Gln Leu Glu Gln Pro Trp Asp Ile Gly
 130 135 140
- Leu Thr Val Asn Asp Ala Phe Arg Pro Val Ser Lys Phe Phe Asp Arg 145 150 155 160
- Val Gln Arg Pro Glu Gln Leu Phe Ser Ile Ala Leu Ala Ala Met Arg 165 170 175
- Val Leu Thr Asp Pro Ala Glu Thr Gly Ala Val Thr Ile Ala Leu Pro 180 185 190
- Glu Asp Val Gln Ala Glu Met Leu Asp Val Pro Val Glu Phe Leu Gln 195 200 205
- Asp Arg Glu Trp His Ile Arg Arg Pro Arg Pro Glu Arg Ala Ala Leu 210 215 220
- Ala Arg Ala Ile Glu Val Ile Lys Asn Ala Lys Asn Pro Met Ile Ile 225 230 235 240
- Ala Gly Gly Gly Val Leu Tyr Ser Asp Ala Glu Thr Gln Leu Gln Ala 245 250 255
- Leu Val Glu Gln Thr Gly Ile Pro Val Gly Thr Ser Gln Ala Gly Gly 260 265 270
- Gly Val Leu Ala Trp Asp His Ala Gln Asn Leu Gly Gly Val Gly Ala 275 280 285
- Thr Gly Thr Leu Ala Ala Asn Arg Ile Ala Gly Asp Ala Asp Val Ile

270 275 act atg atg aac acc ggt gct gtg cac ggt gct gct ctt ggc gca gct 979 Thr Met Met Asn Thr Gly Ala Val His Gly Ala Ala Leu Gly Ala Ala 280 285 gag gtt gca gca acc aag act gag ctt gga ttc gat cct gag gct cac 1027 Glu Val Ala Ala Thr Lys Thr Glu Leu Gly Phe Asp Pro Glu Ala His 295 300 ttc gcg atc gac gat gag gtt atc gct cac acc cgc tcc ctc gca gag 1075 Phe Ala Ile Asp Asp Glu Val Ile Ala His Thr Arg Ser Leu Ala Glu 310 315 320 cgc gct gca cag aag aag gct gca tgg cag gtc aag ttc gat gag tgg 1123 Arg Ala Ala Gln Lys Lys Ala Ala Trp Gln Val Lys Phe Asp Glu Trp 330 335 gca gct gcc aac cct gag aac aag gct ctg ttc gat cgc ctg aac tcc Ala Ala Ala Asn Pro Glu Asn Lys Ala Leu Phe Asp Arg Leu Asn Ser 345 350 cgt gag ctt cca gcg ggc tac gct gac gag ctc cca aca tgg gat gca 1219 Arg Glu Leu Pro Ala Gly Tyr Ala Asp Glu Leu Pro Thr Trp Asp Ala 360 365 gat gag aag ggc gtc gca act cgt aag gct tcc gag gct gca ctt cag 1267 Asp Glu Lys Gly Val Ala Thr Arg Lys Ala Ser Glu Ala Ala Leu Gln 375 380 gea ctg ggc aag acc ctt cct gag ctg tgg ggc ggt tcc gct gac ctc 1315 Ala Leu Gly Lys Thr Leu Pro Glu Leu Trp Gly Gly Ser Ala Asp Leu 390 395 1363 gca ggt tcc aac aac acc gtg atc aag ggc tcc cct tcc ttc ggc cct Ala Gly Ser Asn Asn Thr Val Ile Lys Gly Ser Pro Ser Phe Gly Pro gag too ato too acc gag acc tgg tot gct gag cot tac ggc cgt aac 1411 Glu Ser Ile Ser Thr Glu Thr Trp Ser Ala Glu Pro Tyr Gly Arg Asn 430 ctg cac ttc ggt atc cgt gag cac gct atg gga tcc atc ctc aac ggc 1459 Leu His Phe Gly Ile Arg Glu His Ala Met Gly Ser Ile Leu Asn Gly 1507 att tee etc cae ggt gge ace ege cea tae gge gga ace tte etc ate Ile Ser Leu His Gly Gly Thr Arg Pro Tyr Gly Gly Thr Phe Leu Ile 460 tto too gao tao atg ogt oot goa gtt ogt ott goa got oto atg gag 1555 Phe Ser Asp Tyr Met Arg Pro Ala Val Arg Leu Ala Ala Leu Met Glu 470 ace gae get tae tae gte tgg ace cae gae tee ate ggt etg gge gaa 1603 Thr Asp Ala Tyr Tyr Val Trp Thr His Asp Ser Ile Gly Leu Gly Glu 490 495 gat ggc cca acc cac cag cct gtt gaa acc ttg gct gca ctg cgc gcc

Asp Gly Pro Thr His Gln Pro Val Glu Thr Leu Ala Ala Leu Arg Ala

510

505

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		gct Ala 40														259
		gct Ala														307
		cag Gln														355
tgt Cys	ggc Gly	cac His	tcc Ser	tct Ser 90	ttg Leu	acc Thr	cag Gln	tac Tyr	atc Ile 95	cag Gln	ctt Leu	tac Tyr	ttg Leu	ggt Gly 100	gga Gly	403
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		cca Pro 120														499
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PCT/IB00/00943

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295

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Ala Lys Val	Asp Arg	Pro Ass		t Ile	_	Ile Pro 140	Ala	Thr	Pro	
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Ile Tyr Asp Leu Ala Asn Arg Gly Leu Leu Pro Pro Gly Phe Ser Leu 50 55 60

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Val Trp Glu Arg Leu Ala Glu Gly Met Glu Phe Val Arg Gly Asn Phe 100 105 110

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Asp Lys Thr Arg Gly Thr Ala Gly Asn Trp Ala Tyr Tyr Leu Ser Ile 130 135 140

Pro Pro Asp Ser Phe Thr Ala Val Cys His Gln Leu Glu Arg Ser Gly 145 150 155 160

Met Ala Glu Ser Thr Glu Glu Ala Trp Arg Arg Val Ile Ile Glu Lys 165 170 175

Pro Phe Gly His Asn Leu Glu Ser Ala His Glu Leu Asn Gln Leu Val 180 185 190

Asn Ala Val Phe Pro Glu Ser Ser Val Phe Arg Ile Asp His Tyr Leu 195 200 205

Gly Lys Glu Thr Val Gln Asn Ile Leu Ala Leu Arg Phe Ala Asn Gln 210 215 220

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Thr Met Ala Glu Asp Ile Gly Leu Gly Gly Arg Ala Gly Tyr Tyr Asp 245 250 255

Gly Ile Gly Ala Ala Arg Asp Val Ile Gln Asn His Leu Ile Gln Leu 260 265 270

Leu Ala Leu Val Ala Met Glu Glu Pro Ile Ser Phe Val Pro Ala Gln 275 280 285

Leu Gln Ala Glu Lys Ile Lys Val Leu Ser Ala Thr Lys Pro Cys Tyr 290 295 300

Pro Leu Asp Lys Thr Ser Ala Arg Gly Gln Tyr Ala Ala Gly Trp Gln 305 310 315 320

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Thr Ile Arg Arg Glu Lys Glu Ile Ser Ala Arg Gly Leu His Phe Val 115 120 125

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- Leu Glu Ser Ile Ala Ala Asn Val Asp Gly Thr Pro Cys Val Thr His 165 170 175
- Ile Gly Pro Asp Gly Ala Gly His Phe Val Lys Met Val His Asn Gly 180 185 190
- Ile Glu Tyr Ala Asp Met Gln Val Ile Gly Glu Ala Tyr His Leu Leu 195 200 205
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- Lys Ala Ala Leu Asp Leu Gly Ile Ala Thr Thr Gly Ile Gly Glu Ala 275 280 285
- Val Phe Ala Arg Ala Leu Ser Gly Ala Thr Ser Gln Arg Ala Ala Ala 290 295 300
- Gln Gly Asn Leu Pro Ala Gly Val Leu Thr Asp Leu Glu Ala Leu Gly 305 310 315 320
- Val Asp Lys Ala Gln Phe Val Glu Asp Val Arg Arg Ala Leu Tyr Ala 325 330 335
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- Ser Asp Glu Asn Asn Trp Asp Val Asp Pro Arg Asp Leu Ala Thr Ile 355 360 365
- Trp Arg Gly Gly Cys Ile Ile Arg Ala Lys Phe Leu Asn Arg Ile Val 370 375 380
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ggc Gly	ggc Gly 135	Glu	gaa Glu	ggc	gca Ala	ctc Leu 140	Asn	ggc	cca Pro	tco Ser	Ile 145	atg Met	Pro	ggt Gly	ggc	547
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Trp Thr Ser Gln Gly Ala Gly Glu Gly Leu Val Leu Leu Asp Glu Pro 180 Ser Ser Thr Arg Tyr Pro Ala Ala Pro Gly Gln Asp Glu Val Val Val 200 205 195 Ser Gly Ser Leu Ala Gly Ile Val Arg Tyr Ala Ala Gly Arg Gly Ser Asp Gly Val Thr Ser Ser Thr Gly Glu Val Pro Glu Pro Pro Arg Trp 235 Leu <210> 251 <211> 1575 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1552) <223> RXN00999 <400> 251 cctcctgtga cctggtaaaa tcgccactac ccccaaatgg tcacaccttt taggccgatt 60 ttgctgacac cgggctatgc cgtcaagtac gatcaataac atg act aat gga gat 115 Met Thr Asn Gly Asp aat etc qca cag atc qqc qtt qta qqc cta qca gta atg qqc tca aac 163 Asn Leu Ala Gln Ile Gly Val Val Gly Leu Ala Val Met Gly Ser Asn ctc qcc cgc aac ttc qcc cgc aac ggc aac act gtc gct gtc tac aac 211 Leu Ala Arg Asn Phe Ala Arg Asn Gly Asn Thr Val Ala Val Tyr Asn 25 30 cgc agc act gac aaa acc gac aag ctc atc gcc gat cac ggc tcc gaa 259 Arg Ser Thr Asp Lys Thr Asp Lys Leu Ile Ala Asp His Gly Ser Glu 307 ggc aac ttc atc cct tct gca acc gtc gaa gag ttc gta gca tcc ctg Gly Asn Phe Ile Pro Ser Ala Thr Val Glu Glu Phe Val Ala Ser Leu gaa aag cca cgc cgc gcc atc atc atg gtt cag gct ggt aac gcc acc 355 Glu Lys Pro Arg Arg Ala Ile Ile Met Val Gln Ala Gly Asn Ala Thr gac gca gtc atc aac cag ctg gca gat gcc atg gac gaa ggc gac atc 403 Asp Ala Val Ile Asn Gln Leu Ala Asp Ala Met Asp Glu Gly Asp Ile 90 100 atc atc gac ggc ggc aac gcc ctc tac acc gac acc att cgt cgc gag Ile Ile Asp Gly Gly Asn Ala Leu Tyr Thr Asp Thr Ile Arg Arg Glu 105 110 115

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cgc Arg	tac Tyr	ccc Pro	gcc Ala	gcc Ala 200	cca Pro	ggg Gly	cag Gln	gac Asp	gag Glu 205	gta Val	gta Val	gtg Val	tcc Ser	ggt Gly 210	agc Ser	680
ctt Leu	gca Ala	ggc Gly	att Ile 215	gtt Val	cgc Arg	tac Tyr	gcc Ala	gct Ala 220	ggc Gly	cgc Arg	ggt Gly	tcc Ser	gat Asp 225	gga Gly	gtc Val	728
act Thr	tct Ser	tcc Ser 230	act Thr	gga Gly	gag Glu	gtt Val	cca Pro 235	gag Glu	cca Pro	ccg Pro	cgc Arg	tgg Trp 240	ctg Leu			770
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Phe His Asp Leu Pro Leu Glu Glu Arg Leu Thr Leu Ala Arg Leu Gly
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Thr Ser His Tyr Ser Arg Gln Leu Ser Leu Val Asp Asn Ala Glu Phe
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40 45 50

cac gtg gca tac aac gcc atc gca ctg tgc aac ctc atg cac tgg gca 248 His Val Ala Tyr Asn Ala Ile Ala Leu Cys Asn Leu Met His Trp Ala 55 60 65

aat act ggt gag gaa acc cca atg tac gtg tcg cca gaa gcg cgc aac 296 Asn Thr Gly Glu Glu Thr Pro Met Tyr Val Ser Pro Glu Ala Arg Asn 70 75 80

gag gaa att gcc tac ggt tcc acg ctc aat ccc gat gcg ttg cgt aac 344 Glu Glu Ile Ala Tyr Gly Ser Thr Leu Asn Pro Asp Ala Leu Arg Asn 85 90 95

ctg cat gaa cac tcc gtc gca cgc ctg gac gtg gct tgg cgt gaa acg 392 Leu His Glu His Ser Val Ala Arg Leu Asp Val Ala Trp Arg Glu Thr 100 115

tct gaa gat gct tgg tca cac gag gtt ctg aca gct cag gga cgc act $\ 440$ Ser Glu Asp Ala Trp Ser His Glu Val Leu Thr Ala Gln Gly Arg Thr $\ 120$ $\ 125$ $\ 130$

gtc cca gct agt gaa aca ttg tgg atg cgt tcc cgc gaa gtc tgg atc 488 Val Pro Ala Ser Glu Thr Leu Trp Met Arg Ser Arg Glu Val Trp Ile 135 140 145

cac gca gtt gac ctc ggt gca gtg gca acc ttt ggc gac atc cca gag 536 His Ala Val Asp Leu Gly Ala Val Ala Thr Phe Gly Asp Ile Pro Glu 150 155 160

Asp Arg Leu Asn Ser Arg Glu Leu Pro Ala Gly Tyr Ala Asp Glu Leu 355 360 365

- Pro Thr Trp Asp Ala Asp Glu Lys Gly Val Ala Thr Arg Lys Ala Ser 370 380
- Glu Ala Ala Leu Gln Ala Leu Gly Lys Thr Leu Pro Glu Leu Trp Gly 385 390 395 400
- Gly Ser Ala Asp Leu Ala Gly Ser Asn Asn Thr Val Ile Lys Gly Ser 405 410 415
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- Pro Tyr Gly Arg Asn Leu His Phe Gly Ile Arg Glu His Ala Met Gly 435 440 445
- Ser Ile Leu Asn Gly Ile Ser Leu His Gly Gly Thr Arg Pro Tyr Gly 450 455 460
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- Gly Thr Lys Glu Lys Ala Ala Glu Gly Val Arg Arg Gly Gly Tyr Val 565 570 575
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- Gly Ser Glu Val Gln Leu Ala Val Asn Ala Ala Lys Ala Leu Glu Ala 595 600 605
- Glu Gly Val Ala Ala Arg Val Val Ser Val Pro Cys Met Asp Trp Phe 610 615 620
- Gln Glu Gln Asp Ala Glu Tyr Ile Glu Ser Val Leu Pro Ala Ala Val 625 630 635 640
- Thr Ala Arg Val Ser Val Glu Ala Gly Ile Ala Met Pro Trp Tyr Arg
 645 650 655
- Phe Leu Gly Thr Gln Gly Arg Ala Val Ser Leu Glu His Phe Gly Ala
 660 665
- Ser Ala Asp Tyr Gln Thr Leu Phe Glu Lys Phe Gly Ile Thr Thr Asp

Val Arg Val Leu Ala Ala Asp Ala Val Glu Asn Cys Gly Ser Gly His Pro Gly Thr Ala Met Ser Leu Ala Pro Leu Ala Tyr Thr Leu Tyr Gln 55 Arg Val Met Asn Val Asp Pro Gln Asp Thr Asn Trp Ala Gly Arg Asp Arg Phe Val Leu Ser Cys Gly His Ser Ser Leu Thr Gln Tyr Ile Gln Leu Tyr Leu Gly Gly Phe Gly Leu Glu Met Asp Asp Leu Lys Ala Leu Arg Thr Trp Asp Ser Leu Thr Pro Gly His Pro Glu Tyr Arg His Thr 120 Lys Gly Val Glu Ile Thr Thr Gly Pro Leu Gly Gln Gly Leu Ala Ser 130 135 Ala Val Gly Met Ala Met Ala Ala Arg Arg Glu Arg Gly Leu Phe Asp Pro Thr Ala Ala Glu Gly Glu Ser Pro Phe Asp His His Ile Tyr Val 170 Ile Ala Ser Asp Gly Asp Leu Gln Glu Gly Val Thr Ser Glu Ala Ser 185 Ser Ile Ala Gly Thr Gln Gln Leu Gly Asn Leu Ile Val Phe Trp Asp Asp Asn Arg Ile Ser Ile Glu Asp Asn Thr Glu Ile Ala Phe Asn Glu 215 Asp Val Val Ala Arg Tyr Lys Ala Tyr Gly Trp Gln Thr Ile Glu Val 230 235 Glu Ala Gly Glu Asp Val Ala Ala Ile Glu Ala Ala Val Ala Glu Ala Lys Lys Asp Thr Lys Arg Pro Thr Phe Ile Arg Val Arg Thr Ile Ile 265 Gly Phe Pro Ala Pro Thr Met Met Asn Thr Gly Ala Val His Gly Ala Ala Leu Gly Ala Ala Glu Val Ala Ala Thr Lys Thr Glu Leu Gly Phe Asp Pro Glu Ala His Phe Ala Ile Asp Asp Glu Val Ile Ala His Thr 305 310 315 Arg Ser Leu Ala Glu Arg Ala Ala Gln Lys Lys Ala Ala Trp Gln Val Lys Phe Asp Glu Trp Ala Ala Ala Asn Pro Glu Asn Lys Ala Leu Phe 345

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			ggc tcc gag gtt c Gly Ser Glu Val G 595	-
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Val Glu Ala Gly I.			ttc ttg ggc acc c Phe Leu Gly Thr G 660	
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Asn Tyr Pro Ser As	p Trp Ser A	sp Val Asp Thr 25	Lys Ala Val Asp T	hr

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aac cgt ggc gac gcc cgc gcc tcc gac ctc gtc gca tta gcc aaa gaa Asn Arg Gly Asp Ala Arg Ala Ser Asp Leu Val Ala Leu Ala Lys Glu 330 335 340	1123
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Leu Asp Ser Ser Leu Ala Gln Glu Ile Ala Ala Ile Asp Gly Val Glu 1 5 10 10 15 Leu Asp Ser Glu Val Thr Phe Ala Asp Leu Thr Thr Leu Arg Ile Gly 20 25 30 Gly Lys Pro Arg Ser Ala Val Arg Cys Gln Thr Thr Glu Ala Leu Val 35 40 45 Ser Ala Ile Lys Leu Leu Asp Asp Ala Ser Leu Pro Leu Leu Ile Val	
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Gln Leu Ala	Lys Glu Met 85	Cys Gly Gly	Ser Leu Leu Gly 90	Lys Arg Val 95
Thr Val Leu	Gly Ala Ala 100	Phe Lys Pro 105	Asn Ser Asp Asp	Val Arg Asp 110
Ser Pro Ala 115	Leu Ser Val	Ala Gly Ser 120	Leu Ser Leu Gln 125	Gly Ala Ala
Val Ser Val 130	Tyr Asp Pro	Glu Ala Met 135	Asp Asn Ala Arg 140	Arg Val Phe
Pro Thr Leu 145	Ser Tyr Ala 150	Ser Ser Thr	Lys Glu Ala Leu 155	Ile Asp Ala 160
His Leu Val	Val Leu Ala 165	Thr Glu Trp	Gln Glu Phe Arg 170	Asp Leu Asp 175
Pro Glu Val	Ala Gly Gly 180	Val Val Glu 185	Lys Arg Ala Ile	Ile Asp Gly 190
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			atc ggc gga aaa Ile Gly Gly Lys	

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cac His	ctc Leu	gtc Val	gtt Val	ctt Leu 165	gcc Ala	act Thr	gaa Glu	tgg Trp	caa Gln 170	gaa Glu	ttc Phe	cgc Arg	gac Asp	ctt Leu 175	gac Asp	528
ccc Pro	gaa Glu	gtg Val	gcg Ala 180	gga Gly	ggg Gly	gtc Val	gtc Val	gag Glu 185	aag Lys	cgc Arg	gct Ala	att Ile	att Ile 190	gat Asp	ggc Gly	576
cga Arg	aac Asn	gtc Val 195	ctc Leu	gat Asp	gtt Val	gcc Ala	aaa Lys 200	tgg Trp	aag Lys	gcc Ala	gcc Ala	ggt Gly 205	tgg Trp	gaa Glu	atg Met	624
gaa Glu	gcg Ala 210	ctc Leu	ggc Gly	cgc Arg	aac Asn	ctt Leu 215	tagt	gcgg	gtg d	gatca	aggco	ia ad	ic			668
<211 <212)> 26 l> 21 2> PR 3> Co	5 .T	bact	eriu	ım gl	utan	nicum	l.								
)> 26 Val		Glu	Ile	Cvs	Glu	Pro	Thr	Glv	Ala	Asn	Ala	Val	Δ1 =	I.e.i	
Ţ				5				•	10					15		
Val	Asp	Ala	Ile 20	Gly	His	Asp	Asp	Arg 25	Ile	Gly	Arg	Lys	Phe 30	Leu	Gly	
Ala	Gly	Leu	Gly	Phe	Gly	Gly	Gly	Cys	Leu	Pro	Lys	Asp	Ile	Arg	Ala	

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gcg ggc ctg gga ttc ggt ggc ggt tgt ttg cct aaa gac atc cgc gct

20

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gaa	attt	cgg	caac	gccg	aa t	gtaa	gtta	g tg	tcga	atgc				tcg Ser		115
aat Asn	tac Tyr	gac Asp	tta Leu	atc Ile 10	gtt Val	gta Val	ggc Gly	tcc Ser	ggc Gly 15	ctc Leu	ttc Phe	ggg Gly	ctc Leu	acc Thr 20	gtg Val	163
gct Ala	gag Glu	cgt Arg	gca Ala 25	gct Ala	agc Ser	cag Gln	ctg Leu	ggt Gly 30	aag Lys	aaa Lys	gtc Val	ctc Leu	atc Ile 35	gtt Val	gaa Glu	211
								gct Ala								259
acc Thr	ggc Gly 55	att Ile	gaa Glu	atc Ile	cac His	aaa Lys 60	tac Tyr	ggc Gly	gcg Ala	cac His	ctc Leu 65	ttc Phe	cac His	acc Thr	tcc Ser	307
aac Asn 70	aca Thr	cgc Arg	gtg Val	tgg Trp	gaa Glu 75	tac Tyr	gtc Val	aac Asn	cag Gln	ttc Phe 80	acc Thr	agt Ser	ttc Phe	acc Thr	ggc Gly 85	355
tac Tyr	cag Gln	cac His	cgc Arg	gtc Val 90	ttc Phe	gca Ala	atg Met	cac His	aac Asn 95	ggc Gly	acċ Thr	gcc Ala	tac Tyr	caa Gln 100	ttc Phe	403
								cag Gln 110								451
cca Pro	gat Asp	gaa Glu 120	gcc Ala	cgt Arg	gag Glu	ctc Leu	atc Ile 125	aag Lys	gaa Glu	cag Gln	tct Ser	gca Ala 130	gaa Glu	atc Ile	gat Asp	499
tcc Ser	tcc Ser 135	gac Asp	gcc Ala	acc Thr	aac Asn	ctc Leu 140	gaa Glu	gaa Glu	aag Lys	gcc Ala	att Ile 145	tcc Ser	ctc Leu	att Ile	ggt Gly	547
cgc Arg 150	cca Pro	ctt Leu	tac Tyr	gag Glu	gca Ala 155	ttc Phe	atc	cgc Arg	gac Asp	tac Tyr 160	acc Thr	gca Ala	aag Lys	cag Gln	tgg Trp 165	595
cag Gln																598

gac gac cga gac atg ctg aag cag tac cgc ctt ctg gct gct gaa gag Asp Asp Arg Asp Met Leu Lys Gln Tyr Arg Leu Leu Ala Ala Glu Glu 100 105 110	336
gct gct aat aat aag gtg ctg ttc ggc ggt cga ctg ggc acg tac cag Ala Ala Asn Asn Lys Val Leu Phe Gly Gly Arg Leu Gly Thr Tyr Gln 115 120 125	384
tac ctc gac atg cac atg gct atc ggt tct gcg ctg agc atg ttt gac Tyr Leu Asp Met His Met Ala Ile Gly Ser Ala Leu Ser Met Phe Asp 130 135 140	432
aac aag ctg gtg ccg ttc ttt gaa gaa ggc aca ccg cta gag cag gaa Asn Lys Leu Val Pro Phe Phe Glu Glu Gly Thr Pro Leu Glu Glu Glu 145 150 155 160	480
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Thr Gly Asp Phe Gln Gly Thr Pro Val Met Asn Tyr Asn Asp Ala Asp 35 40 45	
Val Pro Phe Thr Arg Ile His Glu Phe Arg His Phe His Pro Glu Arg 50 55 60	
Asp Asp Ser Tyr Pro Lys Asp Lys Thr Val Ile Met Arg Glu Phe Ser 65 . 70 75 80	
Arg Phe Ala Asp Asn Glu Asp Glu Pro Tyr Tyr Pro Ile Asn Thr Pro 85 90 95	
Asp Asp Arg Asp Met Leu Lys Gln Tyr Arg Leu Leu Ala Ala Glu Glu 100 105 110	
Ala Ala Asn Asn Lys Val Leu Phe Gly Gly Arg Leu Gly Thr Tyr Gln 115 120 125	
Tyr Leu Asp Met His Met Ala Ile Gly Ser Ala Leu Ser Met Phe Asp 130 135 140	
Asn Lys Leu Val Pro Phe Phe Glu Glu Gly Thr Pro Leu Glu Glu Glu 145 150 155 160	
Arg Gly His	

Phe Thr Arg Ile His Glu Phe Arg His Phe His Pro Glu Arg Asp Asp Ser Tyr Pro Lys Asp Lys Thr Val Ile Met Arg Glu Phe Ser Arg Phe 310 Ala Asp Asn Glu Asp Glu Pro Tyr Tyr Pro Ile Asn Thr Pro Asp Asp 330 Arg Asp Met Leu Lys Gln Tyr Arg Leu Leu Ala Ala Glu Glu Ala Ala 345 Asn Asn Lys Val Leu Phe Gly Gly Arg Leu Gly Thr Tyr Gln Tyr Leu 355 360 Asp Met His Met Ala Ile Gly Ser Ala Leu Ser Met Phe Asp Asn Lys Leu Val Pro Phe Phe Glu Glu Gly Thr Pro Leu Glu Gln Glu Arg Gly 395 His <210> 257 <211> 512 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (1)..(489) <223> FRXA02596 <400> 257 cet gtg gtc tac acc ggc cca ctc gac ctc tac ttc aac tac gca gag 48 Pro Val Val Tyr Thr Gly Pro Leu Asp Leu Tyr Phe Asn Tyr Ala Glu ggc aag ctg gga tgg cgc acc ctc gac ttt gaa acc gaa gta gta gaa Gly Lys Leu Gly Trp Arg Thr Leu Asp Phe Glu Thr Glu Val Val Glu acc ggt gac ttc caa gga acc cca gtg atg aac tac aac gat gcg gac Thr Gly Asp Phe Gln Gly Thr Pro Val Met Asn Tyr Asn Asp Ala Asp gta cct ttc acc cgc atc cac gag ttc cgt cac ttc cac cca gag cgt Val Pro Phe Thr Arg Ile His Glu Phe Arg His Phe His Pro Glu Arg 50 55 gat gac agt tac ccc aag gat aag acc gtc atc atg cgc gag ttc tcc Asp Asp Ser Tyr Pro Lys Asp Lys Thr Val Ile Met Arg Glu Phe Ser 65 70 cgt ttc gca gat aac gag gat gag cct tat tac cca atc aac act cca Arg Phe Ala Asp Asn Glu Asp Glu Pro Tyr Tyr Pro Ile Asn Thr Pro 85

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<211> 401

<212> PRT

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Phe Gly Leu Thr Val Ala Glu Arg Ala Ala Ser Gln Leu Gly Lys Lys 20 25 30

Val Leu Ile Val Glu Arg Arg Ser His Leu Gly Gly Asn Ala Tyr Ser 35 40 45

Glu Ala Glu Pro Glu Thr Gly Ile Glu Ile His Lys Tyr Gly Ala His
50 55 60

Leu Phe His Thr Ser Asn Thr Arg Val Trp Glu Tyr Val Asn Gln Phe 65 70 75 80

Thr Ser Phe Thr Gly Tyr Gln His Arg Val Phe Ala Met His Asn Gly 85 90 95

Thr Ala Tyr Gln Phe Pro Met Gly Leu Gly Leu Ile Asn Gln Phe Phe
100 105 110

Gly Lys Tyr Tyr Ser Pro Asp Glu Ala Arg Glu Leu Ile Lys Glu Gln 115 120 125

Ser Ala Glu Ile Asp Ser Ser Asp Ala Thr Asn Leu Glu Glu Lys Ala 130 135 140

Ile Ser Leu Ile Gly Arg Pro Leu Tyr Glu Ala Phe Ile Arg Asp Tyr 145 150 155 160

Thr Ala Lys Gln Trp Gln Thr Asp Pro Lys Asn Leu Pro Ala Gly Asn 165 170 175

Ile Thr Arg Leu Pro Val Arg Tyr Asn Phe Asn Asn Arg Tyr Phe Asn 180 185 190

Asp Thr Tyr Glu Gly Leu Pro Thr Asp Gly Tyr Ala Ala Trp Leu Glu 195 200 205

Lys Met Ala Glu His Glu Leu Ile Asp Val Arg Leu Asp Thr Asp Trp 210 215 220

Phe Asp Val Arg Asp Asp Leu Arg Ala Ser Asn Pro Asp Ala Pro Val 225 230 235 240

Val Tyr Thr Gly Pro Leu Asp Leu Tyr Phe Asn Tyr Ala Glu Gly Lys 245 250 255

Leu Gly Trp Arg Thr Leu Asp Phe Glu Thr Glu Val Val Glu Thr Gly
260 265 270

Asp Phe Gln Gly Thr Pro Val Met Asn Tyr Asn Asp Ala Asp Val Pro 275 280 285

cag Gln	act Thr	gat Asp	cca Pro	aag Lys 170	aac Asn	ctc Leu	cca Pro	gcc Ala	ggc Gly 175	Asn	atc Ile	acc Thr	cgc Arg	ctg Leu 180	cca Pro	643
gtt Val	cgc Arg	tac Tyr	aac Asn 185	Phe	aac Asn	aac Asn	cgc Arg	tat Tyr 190	Phe	aac Asn	gac Asp	acc Thr	tac Tyr 195	gaa Glu	ggc Gly	691
ctt Leu	ccc Pro	aca Thr 200	Asp	ggc Gly	tac Tyr	gcg Ala	gca Ala 205	Trp	ttg Leu	gaa Glu	aag Lys	atg Met 210	gca Ala	gag Glu	cat His	739
gag Glu	ctt Leu 215	atc Ile	gac Asp	gtc Val	cgc Arg	ctc Leu 220	Asp	acc Thr	gac Asp	tgg Trp	ttc Phe 225	gac Asp	gtt Val	cgc Arg	gat Asp	787
gac Asp 230	Leu	cgc Arg	gca Ala	agc Ser	aac Asn 235	ccc Pro	gac Asp	gca Ala	cct Pro	gtg Val 240	Val	tac Tyr	acc Thr	ggc Gly	cca Pro 245	835
ctc Leu	gac Asp	ctc Leu	tac Tyr	ttc Phe 250	aac Asn	tac Tyr	gca Ala	gag Glu	ggc Gly 255	aag Lys	ctg Leu	gga Gly	tgg Trp	cgc Arg 260	acc Thr	883
ctc Leu	gac Asp	ttt Phe	gaa Glu 265	acc Thr	gaa Glu	gta Val	gta Val	gaa Glu 270	acc Thr	ggt Gly	gac Asp	ttc Phe	caa Gln 275	gga Gly	acc Thr	931
cca Pro	gtg Val	atg Met 280	aac Asn	tac Tyr	aac Asn	gat Asp	gcg Ala 285	gac Asp	gta Val	cct Pro	ttc Phe	acc Thr 290	cgc Arg	atc Ile	cac His	979
gag Glu	ttc Phe 295	cgt Arg	cac His	ttc Phe	cac His	cca Pro 300	gag Glu	cgt Arg	gat Asp	gac Asp	agt Ser 305	tac Tyr	ccc Pro	aag Lys	gat Asp	1027
aag Lys 310	acc Thr	gtc Val	atc Ile	atg Met	cgc Arg 315	gag Glu	ttc Phe	tcc Ser	cgt Arg	ttc Phe 320	gca Ala	gat Asp	aac Asn	gag Glu	gat Asp 325	1075
gag Glu	cct Pro	tat Tyr	tac Tyr	cca Pro 330	atc Ile	aac Asn	act Thr	cca Pro	gac Asp 335	gac Asp	cga Arg	gac Asp	atg Met	ctg Leu 340	aag Lys	1123
cag Gln	tac Tyr	cgc Arg	ctt Leu 345	ctg Leu	gct Ala	gct Ala	gaa Glu	gag Glu 350	gct Ala	gct Ala	aat Asn	aat Asn	aag Lys 355	gtg Val	ctg Leu	1171
ttc Phe	ggc Gly	ggt Gly 360	cga Arg	ctg Leu	ggc Gly	acg Thr	tac Tyr 365	cag Gln	tac Tyr	ctc Leu	gac Asp	atg Met 370	cac His	atg Met	gct Ala	1219
atc Ile	ggt Gly 375	tct Ser	gcg Ala	ctg Leu	agc Ser	atg Met 380	ttt Phe	gac Asp	aac Asn	aag Lys	ctg Leu 385	gtg Val	ccg Pro	ttc Phe	ttt Phe	1267
gaa Glu 390	gaa Glu	ggc Gly	aca Thr	ccg Pro	cta Leu 395	gag Glu	cag Gln	gaa Glu	cgc Arg	gga Gly 400	cac His	taaa	agga	ag		1313
ggca	tctc	cc a	ca													1326

Ile Asp Lys Asp Gly Ser Phe His Thr Glu Trp Ser Gly Asp Arg

470

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Ile 145	Met	Pro	Gly	Gly	Pro 150	Ala	Lys	Ser	Tyr	Glu 155	Ser	Leu	Gly	Pro	Leu 160
Leu	Glu	Ser	Ile	Ala 165	Ala	Asn	Val	Asp	Gly 170	Thr	Pro	Cys	Val	Thr 175	His
Ile	Gly	Pro	Asp 180	Gly	Ala	Gly	His	Phe 185	Val	Lys	Met	Val	His 190	Asn	Gly
Ile	Glu	Tyr 195	Ala	Asp	Met	Gln	Val 200	Ile	Gly	Glu	Ala	Tyr 205	His	Leu	Leu
Arg	Tyr 210	Ala	Ala	Gly	Met	Gln 215	Pro	Ala	Glu	Ile	Ala 220	Glu	Val	Phe	Lys
Glu 225	Trp	Asn	Ala	Gly	Asp 230	Leu	Asp	Ser	Tyr	Leu 235	Ile	Glu	Ile	Thr	Ala 240
Glu	Val	Leu	Ser	Gln 245	Val	Asp	Ala	Glu	Thr 250	Gly	Lys	Pro	Leu	Ile 255	Asp
Val	Ile	Val	Asp 260	Ala	Ala	Gly	Gln	Lys 265	Gly	Thr	Gly	Arg	Trp 270	Thr	Val
Lys	Ala	Ala 275	Leu	Asp	Leu	Gly	Ile 280	Ala	Thr	Thr	Gly	Ile 285	Gly	Glu	Ala
Val	Phe 290	Ala	Arg	Ala	Leu	Ser 295	Gly	Ala	Thr	Ser	Gln 300	Arg	Ala	Ala	Ala
Gln 305	Gly	Asn	Leu	Pro	Ala 310	Gly	Val	Leu	Thr	Asp 315	Leu	Glu	Ala	Leu	Gly 320
Val	Asp	Lys	Ala	Gln 325	Phe	Val	Glu	Asp	Val 330	Arg	Arg	Ala	Leu	Tyr 335	Ala
Ser	Lys	Leu	Val 340	Ala	Tyr	Ala	Gln	Gly 345	Phe	Asp	Glu	Ile	Lys 350	Ala	Gly
Ser	Asp	Glu 355	Asn	Asn	Trp	Asp	Val 360	Asp	Pro	Arg	Asp	Leu 365	Ala	Thr	Ile
Trp	Arg 370	Gly	Gly	Cys	Ile	Ile 375	Arg	Ala	Lys	Phe	Leu 380	Asn	Arg	Ile	Val
G1u 385	Ala	Tyr	Asp	Ala	Asn 390	Ala	Glu	Leu	Glu	Ser 395	Leu	Leu	Leu	Asp	Pro 400
Tyr	Phe	Lys	Ser	Glu 405	Leu	Gly	Asp	Leu	Ile 410	Asp	Ser	Trp	Arg	Arg 415	Val
Ile	Val	Thr	Ala 420	Thr	Gln	Leu	Gly	Leu 425	Pro	Ile	Pro	Val	Phe 430	Ala	Ser
Ser	Leu	Ser 435	Tyr	Tyr	Asp	Ser	Leu 440	Arg	Ala	Glu	Arg	Leu 445	Pro	Ala	Ala
Leu	Ile 450	Gln	Gly	Gln	Arg	Asp 455	Phe	Phe	Gly	Ala	His 460	Thr	Tyr	Lys	Arg

atc att co Ile Ile Ar 375														1267
aac gct ga Asn Ala Gl 390														1315
ctc ggc ga Leu Gly As														1363
cag ctt go Gln Leu Gl														1411
gac agc ct Asp Ser Le 44	u Arg													1459
cgc gac tt Arg Asp Ph 455														1507
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ggc Gly	ggc Gly 135	gaa Glu	gaa Glu	ggc Gly	gca Ala	ctc Leu 140	aac Asn	ggc Gly	cca Pro	tcc Ser	atc Ile 145	atg Met	cct Pro	ggt Gly	ggc Gly	547
cca Pro 150	Ala	aag Lys	tcc Ser	tac Tyr	gag Glu 155	tcc Ser	ctc Leu	gga Gly	cca Pro	ctg Leu 160	Leu	gag Glu	tcc Ser	atc Ile	gct Ala 165	595
gcc Ala	aac Asn	gtt Val	gac Asp	ggc Gly 170	acc Thr	cca Pro	tgt Cys	gtc Val	acc Thr 175	cac His	atc Ile	ggc Gly	cca Pro	gac Asp 180	ggc Gly	643
gcc Ala	ggc Gly	cac His	ttc Phe 185	gtc Val	aag Lys	atg Met	gtc Val	cac His 190	aac Asn	ggc Gly	atc Ile	gag Glu	tac Tyr 195	gcc Ala	gac Asp	691
atg Met	cag Gln	gtc Val 200	atc Ile	ggc Gly	gag Glu	gca Ala	tac Tyr 205	cac His	ctt Leu	ctc Leu	cgc Arg	tac Tyr 210	gca Ala	gca Ala	ggc Gly	739
atg Met	cag Gln 215	cca Pro	gct Ala	gaa Glu	atc Ile	gct Ala 220	gag Glu	gtt Val	ttc Phe	aag Lys	gaa Glu 225	tgg Trp	aac Asn	gca Ala	ggc Gly	787
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gtg Val	gat Asp	gct Ala	gaa Glu	acc Thr 250	ggc Gly	aag .Lys	cca Pro	cta Leu	atc Ile 255	gac Asp	gtc Val	atc Ile	gtt Val	gac Asp 260	gct Ala	883
gca Ala	ggt Gly	cag Gln	aag Lys 265	ggc Gly	acc Thr	gga Gly	cgt Arg	tgg Trp 270	acc Thr	gtc Val	aag Lys	gct Ala	gct Ala 275	ctt Leu	gat Asp	931
ctg Leu	ggt Gly	att Ile 280	gct Ala	acc Thr	acc Thr	ggc Gly	atc Ile 285	ggc Gly	gaa Glu	gct Ala	gtt Val	ttc Phe 290	gca Ala	cgt Arg	gca Ala	979
ctc Leu	tcc Ser 295	ggc Gly	gca Ala	acc Thr	agc Ser	cag Gln 300	cgc Arg	gct Ala	gca Ala	gca Ala	cag Gln 305	ggc	aac Asn	cta Leu	cct Pro	1027
gca Ala 310	ggt Gly	gtc Val	ctc Leu	acc Thr	gat Asp 315	ctg Leu	gaa Glu	gca Ala	ctt Leu	ggc Gly 320	gtg Val	gac Asp	aag Lys	gca Ala	cag Gln 325	1075
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tac Tyr	gca Ala	cag Gln	ggc Gly 345	ttc Phe	gac Asp	gag Glu	Ile	aag Lys 350	gct Ala	ggc Gly	tcc Ser	gac Asp	gag Glu 355	aac Asn	aac Asn	1171
tgg Trp	gac Asp	gtt Val 360	gac Asp	cct Pro	cgc Arg	gac Asp	ctc Leu 365	gct Ala	acc Thr	atc Ile	tgg Trp	cgc Arg 370	ggc Gly	ggc	tgc Cys	1219

Leu Glu Ile Glu Tyr Val Val Glu Asp Gln Pro Leu Gly Thr Gly Gly
85 90 95

Gly Ile Arg Asn Val Tyr Asp Lys Leu Arg His Asp Thr Ala Ile Val 100 105 110

Phe Asn Gly Asp Val Leu Ser Gly Ala Asp Leu Asn Ser Ile Leu Asp 115 120 125

Thr His Arg Glu Lys Asp Ala Asp Leu Thr Met His Leu Val Arg Val 130 135 140

Ala Asn Pro Arg Ala Phe Gly Cys Val Pro Thr Asp Glu Asp Gly Arg 145 150 155 160

Val Ser Glu Phe Leu Glu Lys Thr Glu Asp Pro Pro Thr Asp Gln Ile 165 170 175

Asn Ala Gly Cys Tyr Val Phe Lys Lys Glu Leu Ile Glu Gln Ile Pro 180 185 190

Ala Gly Arg Ala Val Ser Val Glu Arg Glu Thr Phe Pro Gln Leu Leu 195 200 205

Glu Glu Gly Lys Arg Val Phe Gly His Val Asp Ala Ser Tyr Trp Arg 210 215 220

Asp Met Gly Thr Pro Ser Asp Phe Val Arg Gly Ser Ala Asp Leu Val 225 230 235 240

Arg Gly Ile Ala Tyr Ser Pro Leu Leu Glu Gly Lys Thr Gly Glu Ser
· 245 250 255

Leu Val Asp Ala Ser Ala Gly Val Arg Asp Gly Val Leu Leu Gly 260 265 270

Gly Thr Val Val Gly Arg Gly Thr Glu Ile Gly Ala Gly Cys Arg Val 275 280 285

Asp Asn Thr Val Ile Phe Asp Gly Val Thr Ile Glu Pro Gly Ala Val . 290 295 300

Ile Glu Asn Ser Ile Ile Ser Ser Gly Ala Arg Ile Gly Ala Asn Ala 305 310 315 320

His Ile Ser Gly Cys Ile Ile Gly Glu Gly Ala Gln Val Gly Ala Arg 325 330 335

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1				5					10					15		
Gly	Gly	Lys	Gly 20	Thr	Arg	Leu .	Arg :	Pro : 25	Leu	Thr	Val	Asn	Thr 30	Pro 1	Lys	
Pro	Met	Leu 35	Pro	Thr	Ala	Gly i	His 1	Pro 1	Phe	Leu	Thr	His:	Leu	Leu A	Ala	
Arg	Ile 50	Lys .	Ala .	Ala	Gly :	Ile 5 55	Thr I	lis V	Val	Val	Leu 60	Gly '	ļhr :	Ser E	Phe	
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Ile Lys His Pro Glu Ala Ser Ile Ile Phe Ser Pro Glu Phe Leu Arg 130 135 140

- Glu Gly Arg Ala Phe Tyr Asp Asn Leu Tyr Pro Ser Arg Val Val 145 150 155 160
- Gly Asp Arg Ser Pro Leu Gly Glu Glu Phe Ala Thr Leu Leu Ala Glu 165 170 175
- Gly Ala Lys Glu Lys Pro Pro Ile Leu Leu Thr Asp Ser Thr Glu Ala 180 185 190
- Glu Ala Ile Lys Leu Phe Ser Asn Thr Tyr Leu Ala Leu Arg Val Ala 195 200 205
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- Lys Gln Ile Ile Glu Gly Val Gly Leu Asp Pro Arg Ile Gly Ser His 225 230 235 240
- Tyr Asn Asn Pro Ser Phe Gly Tyr Gly Gly Tyr Cys Leu Pro Lys Asp 245 250 255
- Thr Lys Gln Leu Leu Ala Asn Tyr Lys Asp Val Pro Gln Asn Leu Ile 260 265 270
- Ser Ala Val Val Gln Ala Asn Lys Thr Arg Lys Asp Phe Ile Ala Glu 275 280 285
- Asp Ile Leu Ser Lys Ser Pro Thr Val Val Gly Ile Tyr Arg Leu Val 290 295 300
- Met Lys Ser Gly Ser Asp Asn Phe Arg Ser Ser Ser Ile Gln Gly Val 305 310 315 320
- Met Lys Arg Ile Lys Ala Lys Gly Ile Glu Ile Val Val Phe Glu Pro 325 330 335
- Asn Leu Gly Glu Glu Thr Phe Tyr Asn Ser Lys Ile Leu Asn Asp Ile 340 345 350
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gcc aag gga Ala Lys Gly												1123
act ttc tac Thr Phe Tyr												1171
tac tgc gac Tyr Cys Asp 360	Ile Ile											1219
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ggg Gly 230	gta Val	ggg Gly	ctc Leu	gat Asp	cca Pro 235	cgt Arg	att Ile	gga Gly	tct Ser	cat His 240	tac Tyr	aat Asn	aat Asn	cct Pro	tca Ser 245	835
ttt Phe	gga Gly	tat Tyr	ggc Gly	gga Gly 250	tat Tyr	tgt Cys	ctt Leu	Pro	aaa Lys 255	gat Asp	acg Thr	aaa Lys	Gln	ctt Leu 260	ctc Leu	883
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Val Pro Lys Glu Leu Leu Pro Val Val Asp Thr Pro Gly Ile Glu Leu 35 40 45

Ile Ala Ala Glu Ala Ala Glu Leu Gly Ala Thr Arg Leu Ala Ile Ile 50 55 60

Thr Ala Pro Asn Lys Ala Gly Val Leu Ala His Phe Glu Arg Ser Ser 65 70 75 80

Glu Leu Glu Glu Thr Leu Met Glu Arg Gly Lys Thr Asp Gln Val Glu
85 90 95

Ile Ile Arg Arg Ala Ala Asp Leu Ile Lys Ala Val Pro Val Thr Gln
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Asp Lys Pro Leu Gly Leu Gly His Ala Val Gly Leu Ala Glu Ser Val 115 120 125

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Met Lys Ile Ala Val

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aat cat aaa gtt gtt gca gtt gac att gat gaa gaa cga gtg aaa cta 211 Asn His Lys Val Val Ala Val Asp Ile Asp Glu Glu Arg Val Lys Leu 25 30 35

Val His Glu Gly Lys Arg His Asp Leu Gly Asn Pro Ala Gly Tyr Ile

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Phe Gly Leu Arg His Ala Glu Tyr Gly Ser Lys Ile His Arg Ala Val 265 270 275

aag gaa ata ctc gct gag ttt gaa tct taaaaaggaa accgccttcc 978 Lys Glu Ile Leu Ala Glu Phe Glu Ser 280 285

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Asp Gly Ile Leu Lys His Phe Glu Glu Phe Pro Glu Leu Glu Ala Thr 50 55 60

Leu Glu Ala Arg Gly Lys Thr Asp Gln Leu Asn Lys Val Arg Ala Ala 65 70 75 80

Arg Glu Leu Ile Ala Thr Val Pro Val Val Gln Glu Lys Pro Leu Gly 85 90 95

Leu Gly His Ala Val Gly Leu Ala Glu Ser Val Leu Asp Asp Glu
100 105 110

Asp Val Val Ala Val Met Leu Pro Asp Asp Leu Val Leu Pro Phe Gly
115 120 125

Val Thr Glu Arg Met Ala Glu Val Arg Ala Lys Phe Gly Gly Ser Val 130 135 140

Leu Ala Ala Ile Glu Val Ala Glu Asp Glu Val Ser Asn Tyr Gly Val 145 150 155 160

Phe Lys Leu Gly Glu Leu Asp Ala Glu Ser Glu Ser Glu Gly Ile Arg 165 170 175

Arg Val Val Gly Met Val Glu Lys Pro Ala Pro Glu Asp Ala Pro Ser 180 185 190

Arg Phe Ala Ala Thr Gly Arg Tyr Leu Leu Asp Arg Ala Ile Phe Asp 195 200 205

Ala Leu Arg Arg Ile Glu Pro Gly Ala Gly Glu Leu Gln Leu Thr 210 215 220

Asp Ala Ile Ala Leu Leu Ile Glu Glu Gly His Pro Val His Ile Val 225 230 235 240

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gtg Val 150	gct Ala	gaa Glu	gat Asp	gaa Glu	gtc Val 155	tca Ser	aat Asn	tac Tyr	gga Gly	gta Val 160	ttt Phe	aag Lys	ctc Leu	ggt Gly	gaa Glu 165	595
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							ctg Leu									787
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275 280 285 acc atc ggc agc cac gtt cgc act ggt tct gac acc atg ttt atc gct Thr Ile Gly Ser His Val Arg Thr Gly Ser Asp Thr Met Phe Ile Ala cca gtg acc gtg ggt gac gga gcg tat tcc gga gcc ggt aca gta att 960 Pro Val Thr Val Gly Asp Gly Ala Tyr Ser Gly Ala Gly Thr Val Ile 310 aaa gac gat gtt ccg cca gga gcc ctt gcc gtg tcc ggc gga cgc caa 1008 Lys Asp Asp Val Pro Pro Gly Ala Leu Ala Val Ser Gly Gly Arg Gln 325 cga aac atc gaa ggc tgg gtg caa aag aag cgc cct gga acc gct gca 1056 Arg Asn Ile Glu Gly Trp Val Gln Lys Lys Arg Pro Gly Thr Ala Ala 345 gca caa gcc gca gaa gcc gcc caa aac gtc cac aac cag gaa ggc 1101 Ala Gln Ala Ala Glu Ala Ala Gln Asn Val His Asn Gln Glu Gly 355 360 taagcaggat cctcatgact gct 1124 <210> 266 <211> 367 <212> PRT <213> Corynebacterium glutamicum <400> 266 Thr Asp His Thr Leu Ser Ala Leu Leu Asp Ala His Val Glu Val Pro 10 Thr Ala Val Thr Val Leu Thr Met Arg Leu Asp Asp Pro Thr Gly Tyr Gly Arg Ile Val Arg Asn Glu Glu Gly Glu Val Thr Ala Ile Val Glu 40 Gln Lys Asp Ala Ser Ala Glu Val Gln Ala Ile Asp Glu Val Asn Ser 55 Gly Val Phe Ala Phe Asp Ala Ala Ile Leu Arg Ser Ala Leu Ala Glu Leu Lys Ser Asp Asn Ala Gln Gly Glu Leu Tyr Leu Thr Asp Val Leu Gly Ile Ala Arg Gly Glu Gly His Pro Val Arg Ala His Thr Ala Ala 105 Asp Ala Arg Glu Leu Ala Gly Val Asn Asp Arg Val Gln Leu Ala Glu 115 Ala Gly Ala Glu Leu Asn Arg Arg Thr Val Ile Ala Ala Met Arg Gly

155

Gly Ala Thr Ile Val Asp Pro Ala Thr Thr Trp Ile Asp Val Glu Val

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										cgt Arg 75						240
										tac Tyr						288
										cgc Arg						336
										cgt Arg						384
										atc Ile						432
										tgg Trp 155						480
										ggc Gly						528
										ggt Gly						576
										gta Val						624
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atc Ile 225	cgc Arg	cca Pro	gga Gly	acc Thr	aca Thr 230	ctg Leu	gga Gly	cca Pro	gaa Glu	ggc Gly 235	aag Lys	ctc Leu	ggt Gly	ggc	ttc Phe 240	720
										ggc Gly						768
										gag Glu						816
										gaa Glu						864

His Gln Val Ser Trp Val Asp Ala Ser Glu Leu Asp Leu Ser Tyr Arg Tyr Ser Asn Leu Lys Phe Thr Asn Arg Ala Val Val Leu Ala Ile Glu 185 Leu Gln Leu Leu Thr Asp Gly Leu Ser Ala Pro Leu Arg Phe Gly Glu 200 Leu Gly Arg Arg Leu Ala Ile Ser Glu Ala Glu Pro His Pro Arg Arg 215 Pro Val Arg Met Val Arg Asp Ala Val Leu Glu Leu Arg Arg Ala Lys Gly Met Val Val Glu His Thr Asp His Asp Thr Trp Ser Ala Gly Ser 250 Phe Phe Thr Asn Pro Ile Val Asp Pro Ala Leu Ala Asp Ala Val Phe Glu Lys Val Gly Glu Pro Thr Met Pro Arg Phe Pro Ala Gly Asp Gly Lys Glu Lys Leu Ser Ala Ala Trp Leu Ile Glu Arg Ala Gly Phe Lys 300 Lys Gly His Pro Gly Ala Gly Ala Lys Ala Ser Leu Ser Thr Lys His 310 315 Thr Leu Ala Leu Thr Asn Arg Gly Asp Ala Arg Ala Ser Asp Leu Val 325 330 Ala Leu Ala Lys Glu Ile Arg Asp Gly Val Leu Glu Thr Phe Gly Val 340 Thr Leu Val Pro Glu Pro Val Trp Ile Gly Ile Ser Ile Asp Asp 360 <210> 265 <211> 1124 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (1)..(1101) <223> RXA01216 acc gac cac act ctg tct gca ctg ctg gat gca cac gtg gaa gtt cca 48 Thr Asp His Thr Leu Ser Ala Leu Leu Asp Ala His Val Glu Val Pro 5 acc gct gtc acc gtg ttg acc atg cgt ctg gat gac ccc acc ggc tac 96 Thr Ala Val Thr Val Leu Thr Met Arg Leu Asp Asp Pro Thr Gly Tyr 25 ggc cgc atc gtg cgc aac gaa gac gaa gtc acc gcc atc gtt gag 144

Ser Val Asn Asp Ala Tyr Leu Gln Gln Gly Ala Leu Thr Val Gln Arg 200 205 210	739												
ctg gac cgt ggc gat gtc tgg tta gat acc ggc aca atc gat tcc atg Leu Asp Arg Gly Asp Val Trp Leu Asp Thr Gly Thr Ile Asp Ser Met 215 220 225	787												
tcc gag gcg tct tcc tat gtt gag gtc ctg caa aaa cgt acc ggc aac Ser Glu Ala Ser Ser Tyr Val Glu Val Leu Gln Lys Arg Thr Gly Asr 230 235 240 245	835												
atc atc gga tcc ccc gaa gtc gct gcg tac cgc gaa ggt ttc atc aca Ile Ile Gly Ser Pro Glu Val Ala Ala Tyr Arg Glu Gly Phe Ile Thr 250 255 260	883												
gct gaa gaa ctc aca gtg ctt ggt gag gaa ctg aag aaa tca ggc tac Ala Glu Glu Leu Thr Val Leu Gly Glu Glu Leu Lys Lys Ser Gly Tyr 265 270 275	931												
gga aac tac ctg ctg aga gct ttg taatttacgg tgtggttgtg gag Gly Asn Tyr Leu Leu Arg Ala Leu 280 285	978												
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Ile Thr Lys Gly Ile Ser Lys Gln Leu Met Pro Ile Tyr Asp Lys Pro													
20 25 30													
20 25 30 Met Val Tyr Tyr Pro Leu Thr Thr Leu Ile Gln Ala Gly Ile Lys Asp													
Met Val Tyr Tyr Pro Leu Thr Thr Leu Ile Gln Ala Gly Ile Lys Asp 35 Ile Leu Ile Ile Thr Thr Pro Glu Asp Ser Ala Ser Phe Glu Arg Leu													
Met Val Tyr Tyr Pro Leu Thr Thr Leu Ile Gln Ala Gly Ile Lys Asp 35 Ile Leu Ile Ile Thr Thr Pro Glu Asp Ser Ala Ser Phe Glu Arg Leu 50 Leu Gly Asp Gly Ser Ser Trp Gly Ile Asn Leu Thr Tyr Ala Val Gln													
Met Val Tyr Tyr Pro Leu Thr Thr Leu Ile Gln Ala Gly Ile Lys Asp 35 Ile Leu Ile Ile Thr Thr Pro Glu Asp Ser Ala Ser Phe Glu Arg Leu 50 Leu Gly Asp Gly Ser Ser Trp Gly Ile Asn Leu Thr Tyr Ala Val Gln 65 Pro Ser Pro Asp Gly Leu Ala Gln Ala Phe Ile Ile Gly Glu Glu Phe													
Met Val Tyr Tyr Pro Leu Thr Thr Leu Ile Gln Ala Gly Ile Lys Asp 35 Ile Leu Ile Ile Thr Thr Pro Glu Asp Ser Ala Ser Phe Glu Arg Leu 50 Leu Gly Asp Gly Ser Ser Trp Gly Ile Asn Leu Thr Tyr Ala Val Gln 65 Pro Ser Pro Asp Gly Leu Ala Gln Ala Phe Ile Ile Gly Glu Glu Phe 95 Ile Gly Asp Asp Asp Asp Val Ala Leu Val Leu Gly Asp Asn Ile Phe Asp													
Met Val Tyr Tyr Pro Leu Thr Thr Leu Ile Gln Ala Gly Ile Lys Asp 45 Ile Leu Ile Ile Thr Thr Pro Glu Asp Ser Ala Ser Phe Glu Arg Leu 60 Leu Gly Asp Gly Ser Ser Trp Gly Ile Asn Leu Thr Tyr Ala Val Gln 65 Pro Ser Pro Asp Gly Leu Ala Gln Ala Phe Ile Ile Gly Glu Glu Phe 95 Ile Gly Asp Asp Asp Asp Val Ala Leu Val Leu Gly Asp Asn Ile Phe Asp 100 Gly Ala Gln Leu Gly His Ala Leu Lys Gln Cys Ser Asn Pro Asp Gly													

<213> Corynebacterium glutamicum

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<213> Corynebacterium glutamicum

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Leu Ala Asp Ser Ile Thr Leu Asp Arg Phe Glu Ala Ser Asp Leu Glu 20 25 30

Val Ser Ser Lys Pro Asp Met Thr Pro Val Ser Asp Ala Asp Leu Ala 35 40 45

Thr Glu Glu Ala Leu Arg Glu Lys Ile Ala Thr Ala Arg Pro Ala Asp 50 55 60

Ser Ile Leu Gly Glu Glu Phe Gly Gly Asp Val Glu Phe Ser Gly Arg
65 70 75 80

Gln Trp Ile Ile Asp Pro Ile Asp Gly Thr Lys Asn Tyr Val Arg Gly
85 90 95

Val Pro Val Trp Ala Thr Leu Ile Ala Leu Leu Asp Asn Gly Lys Pro 100 105 110

Val Ala Gly Val Ile Ser Ala Pro Ala Leu Ala Arg Arg Trp Trp Ala 115 120 125

Ser Glu Gly Ala Gly Ala Trp Arg Thr Phe Asn Gly Ser Ser Pro Arg 130 135 140

Lys Leu Ser Val Ser Gln Val Ser Lys Leu Asp Asp Ala Ser Leu Ser 145 150 155 160

Phe Ser Ser Leu Ser Gly Trp Ala Glu Arg Asp Leu Arg Asp Gln Phe 165 170 175

Val Ser Leu Thr Asp Thr Thr Trp Arg Leu Arg Gly Tyr Gly Asp Phe 180 185 190

Phe Ser Tyr Cys Leu Val Ala Glu Gly Ala Val Asp Ile Ala Ala Glu 195 200 205

Pro Glu Val Ser Leu Trp Asp Leu Ala Pro Leu Ser Ile Leu Val Thr 210 215 220

Glu Ala Gly Gly Lys Phe Thr Ser Leu Ala Gly Val Asp Gly Pro His 225 230 235 . 240

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Asp Arg Leu Lys

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cgc Arg	gag Glu 55	Lys	atc :Ile	gcc Ala	acc Thr	gcc Ala 60	Arg	Pro	gcc Ala	gac Asp	tcc Ser 65	Ile	cto Leu	ggt Gly	gaa Glu	307
gaa Glu 70	ttc Phe	ggt Gly	ggc Gly	gac Asp	gta Val 75	Glu	ttc Phe	agc Ser	Gly	cgc Arg 80	Gln	tgg Trp	ato	atc Ile	gac Asp 85	355
ccc Pro	ạtc Ile	gac Asp	ggc Gly	acc Thr 90	Lys	aac Asn	tac Tyr	gtc Val	cgc Arg 95	Gly	gtc Val	ccc Pro	gta Val	tgg Trp 100	gca Ala	403
acc Thr	ctg Leu	atc Ile	gcg Ala 105	ctg Leu	ctc Leu	gac Asp	aac Asn	ggc Gly 110	aaa Lys	ccc Pro	gtc Val	gca Ala	ggt Gly 115	gtc Val	atc Ile	451
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cag Gln 150	gtg Val	tcc Ser	aag Lys	ctt Leu	gac Asp 155	gac Asp	gcc Ala	tcc Ser	ctc Leu	tcc Ser 160	ttc Phe	tcc Ser	tcc Ser	ctc Leu	tcc Ser 165	595
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acc Thr	acc Thr	tgg Trp	cga Arg 185	ctc Leu	cgc Arg	ggc Gly	tac Tyr	ggc Gly 190	gac Asp	ttc Phe	ttc Phe	tcc Ser	tac Tyr 195	tgc Cys	ctc Leu	691
gtc Val	gcc Ala	gaa Glu 200	ggt Gly	gcc Ala	gtc Val	gat Asp	atc Ile 205	gcc Ala	gct Ala	gaa Glu	cca Pro	gaa Glu 210	gtc Val	agc Ser	ctc Leu	739
rrp .	gat Asp 215	ctt Leu	gct Ala	ccc Pro	ctg Leu	tcc Ser 220	atc Ile	ctg Leu	gtc Val	acc Thr	gaa Glu 225	gcc Ala	gga Gly	gga Gly	aag Lys	787
ttc Phe ' 230	acc Thr	tca Ser	ctg Leu	Ala	ggc Gly 235	gtc Val	gat Asp	gga Gly	Pro	cac His 240	ggt Gly	ggc Gly	gat Asp	gca Ala	gta Val 245	835
gcc a Ala '	acc Thr	aac Asn	Gly	atc Ile 250	ctg Leu	cac His	gat Asp	Glu	acg Thr 255	ctg Leu	gat Asp	cgt Arg	tta Leu	aaa Lys 260		880
aga	ctcc	cg g	gttt	tgct	t gg	t										903

Gln Trp Ile Ile Asp Pro Ile Asp Gly Thr Lys Asn Tyr Val Arg Gly

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			105					110	,				115	5		
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gca Ala	tgg Trp 135	cgc Arg	acc Thr	ttc Phe	aac Asn	ggc Gly 140	agc Ser	tcc Ser	cca Pro	cgc Arg	aaa Lys 145	Leu	tcc Ser	gtg Val	tcc Ser	547
cag Gln 150	gtg Val	tcc Ser	aag Lys	ctt Leu	gac Asp 155	gac Asp	gcc Ala	tcc Ser	ctc Leu	Ser 160	Phe	tcc Ser	tcc Ser	ctc Leu	tcc Ser 165	595
ggc	tgg Trp	gcc Ala	gaa Glu	cga Arg 170	gat Asp	ttg Leu	cgc Arg	gat Asp	cag Gln 175	Phe	gtc Val	tcc Ser	cta Leu	act Thr 180	gat Asp	643
acc Thr	acc Thr	tgg Trp	cga Arg 185	ctc Leu	cgc Arg	ggc	tac Tyr	ggc Gly 190	gac Asp	ttc Phe	ttc Phe	tcc Ser	tac Tyr 195	Cys	ctc Leu	691
gtc Val	gcc Ala	gaa Glu 200	ggt Gly	gcc Ala	gtc Val	gat Asp	atc Ile 205	gcc Ala	gct Ala	gaa Glu	cca Pro	gaa Glu 210	gtc Val	agc Ser	ctc Leu	739
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ttc Phe 230	acc Thr	tca Ser	ctg Leu	gct Ala	ggc Gly 235	gtc Val	gat Asp	gga Gly	cca Pro	cac His 240	ggt Gly	ggc Gly	gat Asp	gca Ala	gta Val 245	835
gcc Ala	acc Thr	aac Asn	ggc Gly	atc Ile 250	ctg Leu	cac His	gat Asp	gag Glu	acg Thr 255	ctg Leu	gat Asp	cgt Arg	tta Leu	aaa Lys 260		880
taga	ctcc	cg g	gttt	tgct	t gg	jt										903
<211 <212)> 27 .> 26 !> PR !> Co	00 T	bact	eriu	um gl	utaπ	ıi cum	ı								
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Leu	Ala	Asp	Ser 20	Ile	Thr	Leu	Asp	Arg 25	Phe	Glu	Ala	Ser	Asp 30	Leu	Glu	
Val	Ser	Ser 35	Lys	Pro	Asp	Met	Thr 40	Pro	Val	Ser	Asp	Ala 45	Asp	Leu	Ala	
Thr	Glu 50	Glu .	Ala	Leu	Arg	Glu 55	Lys	Ile	Ala	Thr	Ala 60	Arg	Pro	Ala	Asp	

Ser Ile Leu Gly Glu Glu Phe Gly Gly Asp Val Glu Phe Ser Gly Arg 65 70 75 80

335 325 330 Ser Asn Asn Val Val Val Glu Glu Gly Ala Thr Val Glu Gly Ala Val 345 340 Leu Met Pro Gly Val Arg Ile Gly Lys Gly Ala Val Val Arg His Ala Ile Leu Asp Lys Asn Val Val Val Arg Asp Gly Glu Leu Ile Gly Val 375 Asp Gln Val Arg Asp Ala Gln Arg Phe Lys Val Ser Ala Gly Gly Val 390 395 385 Val Val Val Gly Lys Asn Gln Val Val 405 <210> 277 <211> 903 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(880) <223> RXN00014 <400> 277 catcaaagtg accgccggcg gcgtcgaatg gtccgttgca ggaaacgcgg aagcagttag 60 tgagatetee gaaactttaa gegeactaga etaacaacae atg age aaa tat gea Met Ser Lys Tyr Ala 1 gac gat tta gcc tta gcc ctc gaa ctt gcc gaa ctt gcc gat tcc atc Asp Asp Leu Ala Leu Ala Leu Glu Leu Ala Glu Leu Ala Asp Ser Ile 10 acc ctc gac cgc ttc gaa gcc tct gac ctg gaa gta tcc tcc aag cca Thr Leu Asp Arg Phe Glu Ala Ser Asp Leu Glu Val Ser Ser Lys Pro 25 30 259 gac atq act ccc qtc aqc qat qcc qac ctq gcg acc gaa gaa gca ctc Asp Met Thr Pro Val Ser Asp Ala Asp Leu Ala Thr Glu Glu Ala Leu 40 cgt gag aaa atc gcc acc gcc cgc ccc gcc gac tcc atc ctc ggt gaa 307 Arq Glu Lys Ile Ala Thr Ala Arq Pro Ala Asp Ser Ile Leu Gly Glu 55 gaa ttc qqt qqc gac gta qaa ttc agc ggc cgc cag tgg atc atc gac 355 Glu Phe Gly Gly Asp Val Glu Phe Ser Gly Arg Gln Trp Ile Ile Asp 70 75 403 ccc atc gac ggc acc aaa aac tac gtc cgc ggc gtc ccc gta tgg gca Pro Ile Asp Gly Thr Lys Asn Tyr Val Arg Gly Val Pro Val Trp Ala 95

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Met Val Lys Gly Val Lys Gly Arg Pro Asn Val Leu Ala Ile Val Leu 1 5 10 15

- Ala Gly Glu Gly Lys Arg Leu Phe Pro Leu Thr Glu Asp Arg Ala 20 25 30
- Lys Pro Ala Val Pro Phe Gly Gly Thr Tyr Arg Leu Ile Asp Phe Val 35 40 45
- Leu Ser Asn Leu Val Asn Ser Gly Phe Leu Lys Ile Ala Val Leu Thr
 50 55 60
- Gln Tyr Lys Ser His Ser Leu Asp Arg His Ile Ser Leu Ser Trp Asn 65 70 75 80
- Val Ser Gly Pro Thr Gly Gln Tyr Ile Ala Ser Val Pro Ala Gln Gln 85 90 95
- Arg Leu Gly Lys Arg Trp Phe Thr Gly Ser Ala Asp Ala Ile Leu Gln
 100 105 110
- Ser Leu Asn Leu Ile Ser Asp Glu Lys Pro Asp Tyr Val Ile Val Phe 115 120 125
- Gly Ala Asp His Val Tyr Arg Met Asp Pro Ser Gln Met Leu Asp Glu 130 135 140
- His Ile Ala Ser Gly Arg Ala Val Ser Val Ala Gly Ile Arg Val Pro 145 150 155 160
- Arg Glu Glu Ala Thr Ala Phe Gly Cys Ile Gln Ser Asp Val Asp Gly 165 170 175
- Asn Ile Thr Glu Phe Leu Glu Lys Pro Ala Asp Pro Pro Gly Thr Pro 180 185 190
- Asp Asp Pro Asp Met Thr Tyr Ala Ser Met Gly Asn Tyr Ile Phe Thr 195 200 205
- Thr Glu Ala Leu Ile Gln Ala Leu Lys Asp Asp Glu Asn Asn Glu Asn 210 215 220
- Ser Asp His Asp Met Gly Gly Asp Ile Ile Pro Tyr Phe Val Ser Arg 225 230 235 240
- Asn Asp Ala His Val Tyr Asp Phe Ser Gly Asn Ile Val Pro Gly Ala 245 250 255
- Thr Glu Arg Asp Lys Gly Tyr Trp Arg Asp Val Gly Thr Ile Asp Ala 260 265 270
- Phe Tyr Glu Cys His Met Asp Leu Ile Ser Val His Pro Ile Phe Asn 275 280 285
- Leu Tyr Asn Ser Glu Trp Pro Ile His Thr Thr Ser Glu Gly Asn Leu 290 295 300
- Pro Pro Ala Lys Phe Val Arg Gly Gly Ile Ala Gln Ser Ser Met Val 305 310 315 320
- Ser Ser Gly Ser Ile Ile Ser Ala Gly Thr Val Arg Asn Ser Val Leu

200 205 210

			gat Asp										787
			ccg Pro 235										835
_			aac Asn		-			-					883
			gtc Val										931
			gtg Val										979
			acc Thr										1027
			gcg Ala 315										1075
	-		gtt Val	-				_			-	-	1123
			acg Thr										1171
			gct Ala										1219
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Lys Ser Pro Gly Asp Phe Ala Thr Glu Val Asp Met Ala Ile Glu Ser 35 40 45

His Met Arg Ser Met Leu Asn Met Met Thr Gly Ile Ala Val Ile Gly 50 55 60

Glu Glu Gly Gly Gly Ala Thr Ser Gly Thr Arg Trp Val Ile Asp Pro 65 70 75 80

Ile Asp Gly Thr Ala Asn Phe Ala Ala Ser Asn Pro Met Ser Ala Ile 85 90 95

Leu Val Ser Leu Leu Val Asp Asp Gln Pro Val Leu Gly Ile Thr Ser 100 105 110

Met Pro Met Leu Gly Lys Arg Leu Thr Ala Phe Glu Gly Ser Pro Leu 115 120 125

Met Ile Asn Gly Glu Pro Gln Glu Pro Leu Gln Glu Gln Ser Ser Leu 130 135 140

Val Ser His Ile Gly Phe Ser Ser Met Ala Ser Pro Arg Asn Thr Ala 145 150 155 160

Phe Pro Val Glu Leu Arg Arg Asp Leu Leu Thr Glu Leu Thr Glu Ser 165 170 175

Tyr Leu Arg Pro Arg Ile Thr Gly Ser Val Gly Val Asp Leu Ala Phe 180 185 190

Thr Ala Gln Gly Ile Phe Gly Ala Cys Val Ser Phe Ser Pro His Val 195 200 205

Trp Asp Asn Ser Ala Gly Val Met Leu Met Arg Ala Ala Gly Ala Gln 210 215 220

Val Thr Asp Thr Glu Gly His Pro Trp Ala Pro Gly Arg Gly Val Val 225 230 235 240

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Glu Tyr Lys 275

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<213> Corynebacterium glutamicum

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aac Asn	ttc Phe	gcg Ala	gcg Ala	tcc Ser 90	Asn	ccg Pro	atg Met	agc Ser	gcg Ala 95	atc Ile	ctg Leu	gtg Val	tct Ser	ttg Leu 100	ctt Leu	403
gtc Val	gac Asp	gac Asp	cag Gln 105	ccc Pro	gtc Val	ctg Leu	ggt Gly	att Ile 110	Thr	tcc Ser	atg Met	ccc Pro	atg Met 115	ctg Leu	ggt Gly	451
aaa Lys	cgc Arg	ctc Leu 120	acc Thr	gct Ala	ttt Phe	gaa Glu	ggt Gly 125	tca Ser	ccg Pro	ctg Leu	atg Met	atc Ile 130	aac Asn	ggt Gly	gaa Glu	499
cct Pro	cag Gln 135	gaa Glu	cca Pro	ttg Leu	caa Gln	gaa Glu 140	caa Gln	tcc Ser	agt Ser	ttg Leu	gta Val 145	tcc Ser	cac His	att Ile	ggt Gly	547
ttt Phe 150	agt Ser	tcc Ser	atg Met	gcc Ala	tcc Ser 155	ccg Pro	cgc Arg	aat Asn	aca Thr	gcg Ala 160	ttt Phe	cct Pro	gtg Val	gag Glu	ttg Leu 165	595
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His	Pro	Ser 195	Arg	Ile	Ala	Glu	Pro 200	Asp	Ile	Gln	Lys	Ala 205	Trp	Met	Ser	
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Ser	Val	Ala	Asp	Trp 245	Asp	Trp	Leu	Pro	Gly 250	Arg	Ala	Leu	Ile	Glu 255	Gly	
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cgaa	agago	jct t	tggd	etgca	ng ta	igaaa	aagct	cgg	gttaa	atac			gct Ala			115
	ttg Leu															163
	cag Gln															211
	gcc Ala															259
	aac Asn 55															307

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	acg Thr 215															787
	gac Asp															835
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1	Asp	-		5					10					15		
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l Glu Gln	Asp	Met Ser 35	Ile 20 Asp	5 Lys Glu	Thr His	Ile Leu	Thr Ala 40	Lys 25 Gln	10 Thr Ala	Phe Leu	Val Val	Ile Tyr 45	Ala 30 Asn	15 His Ala	Asp	
Glu Gln Arg	Asp Asp Leu	Met Ser 35 Ala	Ile 20 Asp Trp	5 Lys Glu Arg	Thr His Met	Ile Leu Arg 55	Thr Ala 40 Glu	Lys 25 Gln Asn	10 Thr Ala Gly	Phe Leu Val	Val Val Asp 60	Ile Tyr 45 Thr	Ala 30 Asn Asp	15 His Ala Tyr	Asp Gly Lys	
Glu Gln Arg Thr 65	Asp Asp Leu 50	Met Ser 35 Ala Val	Ile 20 Asp Trp Ser	5 Lys Glu Arg Asp	Thr His Met Val 70	Ile Leu Arg 55 Val	Thr Ala 40 Glu Thr	Lys 25 Gln Asn Asp	10 Thr Ala Gly Ala	Phe Leu Val Asp 75	Val Val Asp 60 Arg	Ile Tyr 45 Thr	Ala 30 Asn Asp	15 His Ala Tyr Glu	Asp Gly Lys Ala 80	
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Glu Asp Met Ile Lys Thr Ile Thr Lys Thr Phe Val Ile Ala His Asp 20 25 30

- Gln Asp Ser Asp Glu His Leu Ala Gln Ala Leu Val Tyr Asn Ala Gly
 35 40 45
- Arg Leu Ala Trp Arg Met Arg Glu Asn Gly Val Asp Thr Asp Tyr Lys 50 55 60
- Thr Ser Val Ser Asp Val Val Thr Asp Ala Asp Arg Ala Ala Glu Ala 65 70 75 80
- Phe Val Ala Gly Val Leu Glu Ala Leu Arg Pro Glu Asp Gly Val Leu 85 90 95
- Gly Glu Glu Gly Ala Asp Arg Ala Ser Lys Ser Gly Lys Thr Trp Val 100 105 110
- Ile Asp Pro Val Asp Gly Thr Tyr Asn Phe Thr Gln Gly Ser Asp Tyr 115 120 125
- Trp Cys Ser Ala Leu Ala Leu Val Glu Gly Asp Pro Ser Ala Pro Ser 130 140
- Arg Val Leu Phe Gly Ala Val His Arg Pro Ala Met Gly Tyr Thr Trp 145 150 155 160
- Phe Gly Gly Pro Gly Ile Arg Thr Thr Leu Asp Gly Lys Glu Leu Asp 165 170 175
- Leu Leu Val Asp Ala Pro Leu Asn Gln Ile Ser Leu Ala Thr Tyr Ile 180 185 190
- His Pro Ser Arg Ile Ala Glu Pro Asp Ile Gln Lys Ala Trp Met Ser 195 200 205
- Val Ala Thr His Pro Ala Thr Leu Arg Met Phe Gly Ala Gly Ser Ile 210 215 220
- Asp Leu Ala Asn Ile Ala Asp Gly Ser Met Gly Ala Trp Val Gln His 225 230 235 240
- Ser Val Ala Asp Trp Asp Trp Leu Pro Gly Arg Ala Leu Ile Glu Gly 245 250 255
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412

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ctt gaa gcg ttg cgg cct gag gac ggc gtg ctt ggc gag gaa ggc gcg Leu Glu Ala Leu Arg Pro Glu Asp Gly Val Leu Gly Glu Gly Ala

Arg Thr Ala Leu Leu Ile Ala Leu Gly Ala Ile Arg Ser Val Glu Thr qgc gca acc atc aac ctt gct gaa agc atc gag gtt taaccatgac 1121 Gly Ala Thr Ile Asn Leu Ala Glu Ser Ile Glu Val 330 1134 ttttaaactc gca <210> 288 <211> 337 <212> PRT <213> Corynebacterium glutamicum Met Ser Val Lys Leu Ala Leu Ile Gly Ala Gly Arg Ile Gly Ser Asn His Ala Arg Leu Ile Thr Asn His Val Ile Gly Ser Glu Leu Val Ala 20 Val Val Asp Pro Thr Pro Asn Ala Glu Thr Leu Ala Asp Glu Leu Gly Ala Val Ala Phe Ser Asn Pro Asp Asp Val Leu Thr Arg Asp Asp Ile Asp Ala Val Leu Ile Ala Thr Pro Ala Arg Thr His Ala Asp Leu Val Val Lys Ala Ala Ala Gly Lys His Val Phe Val Glu Lys Pro Met Ala Val Thr Leu Glu Asp Ala Asp Arg Ala Ile Asn Ala Ala Arg Glu 105 100 Ala Asn Thr Val Leu Gln Val Gly Phe Asn Arg Arg Phe Ala Ala Gly 120 Phe Ala Ala Arg Ala Arg Ile Asp Ala Gly Asp Ile Gly Thr Pro 130 135 Gln Leu Leu Arg Ser Val Thr Arg Asp Pro Gly Pro Phe Thr Ala Asp 155 150 Pro Asn Lys Ile Pro Gln Trp Thr Ile Phe Leu Glu Thr Leu Ile His Asp Phe Asp Ala Leu Cys Tyr Leu Asn Pro Gly Ala Thr Pro Val Glu 185 Val Thr Ala His Ala Asp Cys Leu Val Val Pro Glu Ala Ala Gly Thr 200 Gly Phe Leu Asp Thr Ala Val Val Thr Val Arg Phe Asp Asn Gly Ala

235

Ile Gly Thr Ala Glu Ala Ser Phe Ser Ala Ala Tyr Gly Tyr Asp Val

230

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Cys	Tyr	Leu	185	Pro	Gly	Ala	acc Thr	Pro 190	Val	Glu	Val	Thr	Ala 195	His	Ala	691
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ASP	tcc Ser 295	atc Ile	cgt Arg	acc Thr	aac Asn	acc Thr 300	cct Pro	tcc Ser	aag Lys	gtt Val	cca Pro 305	ggc Gly	gaa Glu	gct Ala	gca Ala	1027
cgc	acc	gca	cta	ctc	atc	gca	ctc	ggc	gcc	atc	cga	agc	gta	gaa	acc	1075

Tyr Gly Pro Arg Gln Asp Pro His Gly Glu Ala Gly Val Val Ala Ile Phe Ala Leu Arg Leu Leu Gly Gly Leu Asp Thr Lys Val Phe Gly Asp 200 Gly Gly Asn Thr Arg Asp Tyr Val Tyr Val Gly Asp Val Val Arg Ala 215 Phe Tyr Leu Ala Ser Gly Glu Ile Gly Gly Gly Glu Arg Phe Asn Ile Gly Thr Ser Val Glu Thr Ser Asp Arg Gln Leu His Thr Leu Val Ala 250 Thr Ala Ala Gly Ser Lys Asp Asp Pro Glu Tyr Ala Pro Ala Arg Leu Gly Asp Val Pro Arg Ser Ala Leu Ser Phe Gly Lys Ala Lys Glu Val 280 Leu Gly Trp Glu Pro Glu Val Asn Ile Glu Gln Gly Val Ala Lys Thr 290 300 295 Val Glu Tyr Phe Arg Thr His <210> 287 <211> 1134 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1111) <223> RXA01887 <400> 287 catctttaca ggaaacccct tgacggcatc aatgggtggt atctagtatc tactagaacg 60 ttatagtaga acgttctagt aaaacttgga aggatgaaaa atg tca gtc aaa ctt Met Ser Val Lys Leu gcc ctc atc ggt gct gga cgc atc gga tca aat cac gca cgc ctg atc Ala Leu Ile Gly Ala Gly Arg Ile Gly Ser Asn His Ala Arg Leu Ile 10 aca aac cac gtg atc ggc tct gaa ctg gtc gcc gtc gtt gac cca act Thr Asn His Val Ile Gly Ser Glu Leu Val Ala Val Val Asp Pro Thr 25 ccc aac gca gaa acc ctc gct gat gaa ttg ggc gcc gtt gcg ttc tct Pro Asn Ala Glu Thr Leu Ala Asp Glu Leu Gly Ala Val Ala Phe Ser 40 aac cca gat gac gtc ctg acc cgc gat gac att gac gcg gtt ttg att 307 Asn Pro Asp Asp Val Leu Thr Arg Asp Asp Ile Asp Ala Val Leu Ile 60

aco Thi	c tct r Ser	gad Ksp	cgo Aro	C Caq Glr 250	ı Lev	cac His	acc Thi	c cto	c gtg 2 Val 255	L Ala	c act	gc Ala	g gca a Ala	a ggt a Gly 260	tcc y Ser)	883
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agt Ser	gca Ala	Leu 280	ı Ser	tto Phe	ggc Gly	aag Lys	gcc Ala 285	Lys	a gaç s Glu	gto Val	g ctt Leu	ggt Gl ₃ 290	/ Trp	g gaç o Glu	g cct Pro	979
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Ile	Asp	Val	Arg	His 85	Ser	Val	Val	Asp	Pro 90	Leu	His	Asp	Ala	Glu 95	Thr	
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Pro	Ser 130	Glu	Phe	Pro	Val	Asp 135	Glu	Thr	Val	Pro	Val 140	Asp	Pro	His	Ser	
Pro 145	Tyr	Ala	Ala	Ser	Lys 150	Val	Ser	Gly	Glu	Ile 155	Tyr	Leu	Asn	Thr	Phe 160	
Arg	His	Leu	Tyr	Gly 165	Leu	Asp	Cys	Ser	His 170	Ile	Ala	Pro	Ala	Asn 175	Val	

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240

Gly Glu Ile Gly Gly Glu Arg Phe Asn Ile Gly Thr Ser Val Glu

235

230

Arg Glu Arg Thr Leu Leu Glu Arg Ser Leu Gln Ala Met Leu Thr Ser 35 40 45

Glu Ser Val Asp Glu Ile Ile Ile Leu Val Ser Pro Asp Met Glu Thr
50 55 60

Tyr Ala Arg Asp Leu Leu Arg Lys Arg Gly Leu Leu Asn Asp Pro Glu 65 70 75 80

Gly Val Arg Val Arg Leu Val His Gly Gly Glu Arg Ala Asp Ser 85 90 95

Val Trp Ala Gly Leu Gln Ala Ile Ser Leu Asp Asp Ala Thr Pro Asp 100 105 110

Ala Ile Val Leu Ile His Asp Ser Ala Arg Ala Leu Thr Pro Pro Gly 115 120 125

Met Ile Ala Arg Val Val Arg Lys Val His Glu Gly Ala Thr Ala Val 130 135 140

Ile Pro Val Leu Pro Val Ser Asp Thr Ile Lys Arg Val Ser Pro Asp 145 150 155 160

Gly Gly Val Val Asp Thr Pro Asn Arg Ala Glu Leu Arg Ala Val 165 170 175

Gln Thr Pro Gln Gly Phe Leu Leu Ser Glu Leu Val Ala Ala Asn Glu 180 185 190

Lys Phe Phe Ala Asp Pro Asn Pro Gly Phe Ile Pro Thr Asp Asp Ala 195 200 205

Ser Leu Met Glu Trp Tyr Gly Ala Asp Val Val Cys Val Gln Gly Asp 210 215 220

Pro Met Ala Phe Lys Val Thr Thr Pro Ile Asp Met Met Leu Ala Gln 225 230 235 240

Arg Ile Thr Asp Glu Ala Glu Pro Thr Ile Phe Glu Val Pro Gly Asp 245 250 255

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70	75	8	0	85
ctc gtg cac g Leu Val His G	gc ggc ggg ga ly Gly Gly Gl 90	g cgc gcg gac to u Arg Ala Asp Se 95	g gtc tgg gca ggc r Val Trp Ala Gly 100	ctt 403 Leu
Gln Ala Ile S			et gca att gtc tta op Ala Ile Val Leu 115	
			c atg att gcg cgc y Met Ile Ala Arg 130	
		y Ala Thr Ala Va	c atc cca gta ctg l Ile Pro Val Leu 145	
			t ggc gga gta gtt p Gly Gly Val Val 0	
			c caa acc cca caa l Gln Thr Pro Gln 180	
Phe Leu Leu S	•		g aaa ttc ttc gcc u Lys Phe Phe Ala 195	-
-	_		c agc ttg atg gaa a Ser Leu Met Glu 210	
		s Val Gln Gly As	c cca atg gcg ttt p Pro Met Ala Phe 225	
Val Thr Thr P	ro Ile Asp Me		a cgc atc acc gac n Arg Ile Thr Asp 0	
		g gta cca ggt ga u Val Pro Gly As 255	c taacccaatc atccc p	ccgcg 888
tag				891
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	rg Leu Gly Gl 20	y Pro Ile Pro Ly 25	s Ala Phe Val Thr 30	Leu

Thr Ala Pro Lys Ser Asn Phe Ala Val Val Gly Leu Tyr Phe Tyr Asp Asn Arg Val Val Asp Ile Ala Lys Ser Ile Lys Pro Ser Ser Arg Gly 185 Glu Leu Glu Ile Thr Ser Val Asn Asp Ala Tyr Leu Gln Gln Gly Ala 200 Leu Thr Val Gln Arg Leu Asp Arg Gly Asp Val Trp Leu Asp Thr Gly 215 Thr Ile Asp Ser Met Ser Glu Ala Ser Ser Tyr Val Glu Val Leu Gln 230 Lys Arg Thr Gly Asn Ile Ile Gly Ser Pro Glu Val Ala Ala Tyr Arg 250 Glu Gly Phe Ile Thr Ala Glu Glu Leu Thr Val Leu Gly Glu Glu Leu Lys Lys Ser Gly Tyr Gly Asn Tyr Leu Leu Arg Ala Leu 280 <210> 283 <211> 891 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(868) <223> RXA02666 <400> 283 geteggegae gaggaagaga agaaggaege attegaegae ttegaegatt eegaegtgga 60 tettgaegat etgagetteg acgaegaaga ttagaegeee atg teg tet aca ega Met Ser Ser Thr Arg atc ccc gtc atc gca ctc ctc gcg gcg gcg ggg cgc gga acc cgc ctc 163 Ile Pro Val Ile Ala Leu Leu Ala Ala Ala Gly Arg Gly Thr Arg Leu 10 ggc gga ccc atc ccc aaa gca ttc gtc acg ttg cgt gaa cgc aca ctt Gly Gly Pro Ile Pro Lys Ala Phe Val Thr Leu Arg Glu Arg Thr Leu 25 tta gag cgc tcg ctc caa gcc atg ctc acc tcc gaa agc gtc gac gaa 259 Leu Glu Arg Ser Leu Gln Ala Met Leu Thr Ser Glu Ser Val Asp Glu 40 45 atc atc atc ctc gtc agc ccc gac atg gaa acc tac gcc cgc gat ttg 307 Ile Ile Ile Leu Val Ser Pro Asp Met Glu Thr Tyr Ala Arg Asp Leu 55 60 ctg cgc aaa cgc ggt ctt ttg aac gac ccc gaa ggg gta cgc gta cgg 355 Leu Arg Lys Arg Gly Leu Leu Asn Asp Pro Glu Gly Val Arg Val Arg

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<212> PRT

<213> Corynebacterium glutamicum

<400> 296

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Ile Ala Ala Asp Ala Val Asp Ala Val Leu Ile Ala Val Pro Gly Gln
35 40 45

Phe His Glu Pro Val Leu Val Pro Ala Leu Glu Ala Gly Leu Pro Ile 50 55 60

Leu Cys Glu Lys Pro Leu Thr Pro Asp Ser Glu Ser Ser Leu Arg Ile
65 70 75 80

Val Glu Leu Glu Gln Lys Leu Asp Lys Pro His Ile Gln Val Gly Phe
85 90 95

Met Arg Arg Phe Asp Pro Glu Tyr Asn Asn Leu Arg Lys Leu Val Glu
100 105 110

Ser Gly Glu Ala Gly Glu Leu Leu Met Leu Arg Gly Leu His Arg Asn 115 120 125

Pro Ser Val Gly Glu Ser Tyr Thr Gln Ser Met Leu Ile Thr Asp Ser 130 135 140

Val Val His Glu Phe Asp Val Ile Pro Trp Leu Ala Gly Ser Arg Val 145 150 155 160

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Gly Leu Lys Glu Pro Ile Leu Val Ile Met Glu Leu Glu Asn Gly Val 180 185 190

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<211> 549

<212> DNA

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ttc act cgc att gaa gat gct atc gca gcc gat gct gtc gac gca gtg 96
Phe Thr Arg Ile Glu Asp Ala Ile Ala Ala Asp Ala Val Asp Ala Val
20 25 30

ctg atc gcc gta cca ggt cag ttc cat gag cca gta ctt gtc cca gca 144 Leu Ile Ala Val Pro Gly Gln Phe His Glu Pro Val Leu Val Pro Ala

45 40 35 cta gaa gca ggc ctt ccc atc ctg tgt gaa aag cca ctg acc cca gat Leu Glu Ala Gly Leu Pro Ile Leu Cys Glu Lys Pro Leu Thr Pro Asp 55 tot gaa too toa otg ogo ato gto gag otg gag cag aag otg gac aag Ser Glu Ser Ser Leu Arg Ile Val Glu Leu Glu Gln Lys Leu Asp Lys cca cac atc cag gtt ggt ttc atg cgc cgc ttc gac cct gag tac aac Pro His Ile Gln Val Gly Phe Met Arg Arg Phe Asp Pro Glu Tyr Asn 85 aac ttg cgc aaa ttg gtg gaa tcc ggc gaa gct ggc gaa ctg ctc atg 336 Asn Leu Arg Lys Leu Val Glu Ser Gly Glu Ala Gly Glu Leu Leu Met 100 ctc cgc ggc ctg cac cgc aac cca agt gtt ggt gag agc tac acc cag Leu Arg Gly Leu His Arg Asn Pro Ser Val Gly Glu Ser Tyr Thr Gln 125 115 120 tee atg etg ate ace gae tee gte gte cae gaa tte gat gte ate eea Ser Met Leu Ile Thr Asp Ser Val Val His Glu Phe Asp Val Ile Pro 130 135 tgg etc gca ggc tcc cga gtt gtc tcc gtt gaa gtg aag tac cca aag Trp Leu Ala Gly Ser Arg Val Val Ser Val Glu Val Lys Tyr Pro Lys 160 145 150 acc tcc tca ctg gcg cac tcc ggc ctc aag gaa cca atc ctg gtg atc Thr Ser Ser Leu Ala His Ser Gly Leu Lys Glu Pro Ile Leu Val Ile 549 atg gag ctc gaa aac ggc gtg Met Glu Leu Glu Asn Gly Val 180 <210> 298 <211> 183 <212> PRT <213> Corynebacterium glutamicum <400> 298

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Leu	Arg	Gly 115	Leu	His	Arg	Asn	Pro 120		Val	Gly	Glu	Ser 125		Thr	Gln	
Ser	Met 130	Leu	Ile	Thr	Asp	Ser 135	Val	Val	His	Glu	Phe 140	Asp	Val	Ile	Pro	
Trp 145	Leu	Ala	Gly	Ser	Arg 150	Val	Val	Ser	Val	Glu 155	Val	Lys	Tyr	Pro	Lys 160	
Thr	Ser	Ser	Leu	Ala 165	His	Ser	Gly	Leu	Lys 170		Pro	Ile	Leu	Val 175	Ile	
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cccc	ggctg	gca a	ggc	gato	et tt	gaaa	aggci atc	t ga: ggt	aaaaa cac	actc gtc	atg Met 1	act Thr	ctt Leu	cgt	atc Ile 5	
gcc Ala	ggcto cctco ctt Leu gca	gca acac ttc Phe	ggc Gly cct	gct Ala 10	ggc Gly	cgc Arg	aggci atc Ile	ggt Gly gtt	cac His 15	gtc Val	atg Met 1 cac His	act Thr gct Ala	ctt Leu gcc Ala	cgt Arg aac Asn	atc Ile 5 att Ile	115
gcc Ala gct Ala	ctt Leu gca Ala	ttc Phe aac Asn	ggc Gly cct Pro 25	gct Ala 10 gat Asp	ggc Gly ctt Leu	cgc Arg gaa Glu gca	atc Ile ctc Leu	ggt Gly gtt Val 30	cac His 15 gtt Val	gtc Val atc Ile	atg Met 1 cac His gcc Ala	act Thr gct Ala gat Asp	ctt Leu gcc Ala cct Pro 35	cgt Arg aac Asn 20	atc Ile 5 att Ile att Ile	115
gcc Ala gct Ala Glu	ctt Leu gca Ala ggc Gly	ttc Phe aac Asn gca Ala 40	ggc Gly cct Pro 25 cag Gln	gct Ala 10 gat Asp cgt Arg	ggc Gly ctt Leu ttg Leu	cgaaa cgc Arg gaa Glu gca Ala	atc Ile ctc Leu gaa Glu 45	ggt Gly gtt Val 30 gcc Ala	cac His 15 gtt Val aat Asn	gtc Val atc Ile ggg Gly	atg Met 1 cac His gcc Ala gca Ala	act Thr gct Ala gat Asp gaa Glu 50	ctt Leu gcc Ala cct Pro 35 gcg Ala	cgt Arg aac Asn 20 ttc Phe	atc Ile 5 att Ile att Ile gca Ala	115 163 211
gcc Ala gct Ala Glu tca Ser	ctt Leu gca Ala ggc Gly cca Pro 55	ttc Phe aac Asn gca Ala 40 gat Asp	ggc Gly cct Pro 25 cag Gln gag Glu	gct Ala 10 gat Asp cgt Arg gtg Val	ggc Gly ctt Leu ttg Leu ttc Phe	cgaaa cgc Arg gaa Glu gca Ala gcc Ala 60 cac	atc Ile ctc Leu gaa Glu 45 cgc Arg	ggt Gly gtt Val 30 gcc Ala gat Asp	cac His 15 gtt Val aat Asn gat	gtc Val atc Ile ggg Gly atc	atg Met 1 cac His gcc Ala gca Ala gat Asp 65 acc	act Thr gct Ala gat Asp gaa Glu 50 ggc Gly	ctt Leu gcc Ala cct Pro 35 gcg Ala atc	cgt Arg aac Asn 20 ttc Phe gtt Val	atc Ile 5 att Ile att Ile gca Ala atc Ile gaa	115163211259

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					cga Arg											499
					cag Gln											547
					gcg Ala 155											595
					gat Asp											643
					atc Ile											691
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acg Thr	ctt Leu 215	cgt Arg	ggc Gly	tca Ser	aag Lys	ggc Gly 220	gag Glu	ttg Leu	atc Ile	aac Asn	atc Ile 225	gtg Val	aac Asn	tcc Ser	cgc Arg	787
cac His 230	tgc Cys	tcc Ser	tac Tyr	ggc Gly	tac Tyr 235	gac Asp	cag Gln	cga Arg	ctt Leu	gag Glu 240	gct Ala	ttc Phe	ggc Gly	tct Ser	aag Lys 245	835
					gac Asp											883
aat Asn	gcg Ala	gaa Glu	agc Ser 265	acc Thr	gag Glu	cag Gln	gca Ala	gat Asp 270	ccg Pro	att Ile	ttċ Phe	aac Asn	ttc Phe 275	ttc Phe	ctc Leu	931
					gct Ala											979
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<211> 335

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<213> Corynebacterium glutamicum

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20 25 30

Ala Asp Pro Phe Ile Glu Gly Ala Gln Arg Leu Ala Glu Ala Asn Gly 35 40 45

Ala Glu Ala Val Ala Ser Pro Asp Glu Val Phe Ala Arg Asp Asp Ile 50 55 60

Asp Gly Ile Val Ile Gly Ser Pro Thr Ser Thr His Val Asp Leu Ile 65 70 75 80

Thr Arg Ala Val Glu Arg Gly Ile Pro Ala Leu Cys Glu Lys Pro Ile 85 90 95

Asp Leu Asp Ile Glu Met Val Arg Ala Cys Lys Glu Lys Ile Gly Asp 100 105 110

Gly Ala Ser Lys Val Met Leu Gly Phe Asn Arg Arg Phe Asp Pro Ser 115 120 125

Phe Ala Ala Ile Asn Ala Arg Val Ala Asn Gln Glu Ile Gly Asn Leu 130 135 140

Glu Gln Leu Val Ile Ile Ser Arg Asp Pro Ala Pro Ala Pro Lys Asp 145 150 155 160

Tyr Ile Ala Gly Ser Gly Gly Ile Phe Arg Asp Met Thr Ile His Asp 165 170 175

Leu Asp Met Ala Arg Phe Phe Val Pro Asn Ile Val Glu Val Thr Ala 180 185 190

Thr Gly Ala Asn Val Phe Ser Gln Glu Ile Ala Glu Phe Asn Asp Tyr 195 200 205

Asp Gln Val Ile Val Thr Leu Arg Gly Ser Lys Gly Glu Leu Ile Asn 210 215 220

Ile Val Asn Ser Arg His Cys Ser Tyr Gly Tyr Asp Gln Arg Leu Glu 225 230 235 240

Ala Phe Gly Ser Lys Gly Met Leu Ala Ala Asp Asn Ile Arg Pro Thr 245 250 255

Thr Val Arg Lys His Asn Ala Glu Ser Thr Glu Gln Ala Asp Pro Ile
260 265 270

Phe Asn Phe Phe Leu Glu Arg Tyr Asp Ala Ala Tyr Lys Ala Glu Leu

285 275 280 Ala Thr Phe Ala Gln Gly Ile Arg Asp Gly Gln Gly Phe Ser Pro Asn 295 Phe Glu Asp Gly Val Ile Ala Leu Glu Leu Ala Asn Ala Cys Leu Glu 315 310 Ser Ala Gln Thr Gly Arg Thr Val Thr Leu Asn Pro Ala Asn Val 325 330 <210> 301 <211> 1206 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1183) <223> RXA01633 <400> 301 gcgaatgcat gccttgaatc agctcaaacc ggccgcaccg tcaccctcaa ccctgccaac 60 gtttagtcaa cgtctagtta atgcctaagg agaaaacctc atg aaa aac atc acc Met Lys Asn Ile Thr atc gga atg gtc ggc gtc ggc cgc att ggc cgc atg cac gtc gcc aac 163 Ile Gly Met Val Gly Val Gly Arg Ile Gly Arg Met His Val Ala Asn 10 atg ctt gcc gtt gct gaa act ttg aag gaa cgc gac ctc aac att gag Met Leu Ala Val Ala Glu Thr Leu Lys Glu Arg Asp Leu Asn Ile Glu 25 atc gtg ctc gca gac gca atg ccc ggt ttt gcg gag cag gtg ggc gcg 259 Ile Val Leu Ala Asp Ala Met Pro Gly Phe Ala Glu Gln Val Gly Ala 40 45 gac atg ggc gtg aag gcg gca agc gtc gat aag ctt att gag gac 307 Asp Met Gly Val Lys Ala Ala Ala Ser Val Asp Lys Leu Ile Glu Asp 65 55 60 ggg gtg gat gcc ctt ttc att gcc acc agc acc gct ggc cac gtc gat 355 Gly Val Asp Ala Leu Phe Ile Ala Thr Ser Thr Ala Gly His Val Asp 70 gtt ttg cgc aag ggc atc gcg gca aag ctg ccg atg ttc tgc gag aag 403 Val Leu Arg Lys Gly Ile Ala Ala Lys Leu Pro Met Phe Cys Glu Lys 100 90 ccg atc gcg tcg gat gtg cct gag tcg ctg aac atc atc cgc gaa att 451 Pro Ile Ala Ser Asp Val Pro Glu Ser Leu Asn Ile Ile Arg Glu Ile 115 105 110 gat gcg gct ggc gcg acg gtt cag gtc ggc cac cag cgc cgt ttt gac

Asp Ala Ala Gly Ala Thr Val Gln Val Gly His Gln Arg Arg Phe Asp

120

ct Le	e gg u Gl; 13	у ту	c ca r Gl	g gaa n Glu	a gct u Ala	aaa Lys 140	: Arg	cgc Arg	cta Lei	a gat ı Asp	gca Ala 145	Gl3	gad Asp	cto Le	ggc Gly	547
tgg Tr _I 150	у пе	t car u Hi:	t tc s Se:	g cto r Lei	c aag 1 Lys 155	Ala	gta Val	tcg Ser	ago Sei	gat Asp 160	Ala	ttt Phe	ccg Pro	g cca Pro	ccg Pro 165	595
gto Va]	g too L Sei	tac Ty	c tgo	c gct s Ala 170	Thr	tct Ser	ggt	gga Gly	ctt Let 175	2 Phe	cgc Arg	gat Asp	gtg Val	tcg Ser 180	ctg Leu	643
Cac His	gat Asp	tto Phe	gad Asp 185) TTE	att : Ile	cgc Arg	tgg Trp	ctg Leu 190	acc Thr	ggc Gly	cag Gln	gat Asp	att Ile 195	Val	gag Glu	691
gtg Val	tac Tyr	gco Ala 200	і гуз	ggc Gly	agc Ser	aac Asn	aac Asn 205	ggc Gly	gac Asp	cca Pro	gaa Glu	atc Ile 210	ggc Gly	gca Ala	gtc Val	739
ggt Gly	gac Asp 215	, тте	gat Asp	acc Thr	gga Gly	gcg Ala 220	gcc Ala	cta Leu	ctc Leu	acg Thr	ctt Leu 225	gcc Ala	gac Asp	ggc	acc	787
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cgc Arg	ctc Leu	gat Asp	gtt Val	atg Met 250	ggc Gly	tct Ser	aaa Lys	gat Asp	tcc Ser 255	acg Thr	atc Ile	gtt Val	ggc Gly	ctg Leu 260	gat Asp	883
gaa Glu	aag Lys	tct Ser	gcg Ala 265	ttc Phe	gct Ala	tct Ser	gcg Ala	gag Glu 270	gag Glu	ggc Gly	atc Ile	gat Asp	ttc Phe 275	cca Pro	acc Thr	931
ggc Gly	gaa Glụ	tcg Ser 280	cac His	cca Pro	acg Thr	ttt Phe	gcc Ala 285	gag Glu	cgc Arg	ttc Phe	gcc Ala	gac Asp 290	gca Ala	tac Tyr	aag Lys	979
aat Asn	gag Glu 295	tgc Cys	att Ile	gcg Ala	ttc Phe	gtg Val 300	gag Glu	ttg Leu	atc Ile	ctg Leu	gga Gly 305	gag Glu	cgg Arg	gaa Glu	aac Asn	1027
cct Pro 310	tgt Cys	acc Thr	cct Pro	gca Ala	gac Asp 315	gct Ala	gtg Val	gct Ala	gcg Ala	gcg Ala 320	att Ile	gtt Val	gcc Ala	gat Asp	gca Ala 325	1075
gct Ala	cag Gln	ctg Leu	tcg Ser	ctg Leu 330	gtc Val	act Thr	ggc (Gly (Glu	cca Pro 335	gtg Val	aag Lys	att Ile	cct Pro	act Thr 340	gta Val	1123
cgg Arg	gaa Glu	att Ile	ctt Leu 345	gaa Glu	ggt Gly	tct Ser	Ala (cag Gln 350	cca Pro	gtt Val	gag Glu	Val .	cgt Arg 355	gcg Ala	ctg Leu	1171
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<211> 361

<212> PRT

<213> Corynebacterium glutamicum

<400> 302

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20 25 30

Asp Leu Asn Ile Glu Ile Val Leu Ala Asp Ala Met Pro Gly Phe Ala 35 40 45

Glu Gln Val Gly Ala Asp Met Gly Val Lys Ala Ala Ala Ser Val Asp
50 55 60

Lys Leu Ile Glu Asp Gly Val Asp Ala Leu Phe Ile Ala Thr Ser Thr 65 70 75 80

Ala Gly His Val Asp Val Leu Arg Lys Gly Ile Ala Ala Lys Leu Pro 85 90 95

Met Phe Cys Glu Lys Pro Ile Ala Ser Asp Val Pro Glu Ser Leu Asn 100 105 110

Ile Ile Arg Glu Ile Asp Ala Ala Gly Ala Thr Val Gln Val Gly His 115 120 125

Gln Arg Arg Phe Asp Leu Gly Tyr Gln Glu Ala Lys Arg Arg Leu Asp 130 135 140

Ala Gly Asp Leu Gly Trp Leu His Ser Leu Lys Ala Val Ser Ser Asp 145 150 155 160

Ala Phe Pro Pro Pro Val Ser Tyr Cys Ala Thr Ser Gly Gly Leu Phe 165 170 175

Arg Asp Val Ser Leu His Asp Phe Asp Ile Ile Arg Trp Leu Thr Gly 180 185 190

Gln Asp Ile Val Glu Val Tyr Ala Lys Gly Ser Asn Asn Gly Asp Pro 195 200 205

Glu Ile Gly Ala Val Gly Asp Ile Asp Thr Gly Ala Ala Leu Leu Thr 210 215 220

Leu Ala Asp Gly Thr Leu Ala Thr Ala Ile Ala Thr Arg Tyr Asn Gly 225 230 235 240

Ala Gly His Asp Val Arg Leu Asp Val Met Gly Ser Lys Asp Ser Thr

Ile Val Gly Leu Asp Glu Lys Ser Ala Phe Ala Ser Ala Glu Glu Gly 260 265 270

Ile Asp Phe Pro Thr Gly Glu Ser His Pro Thr Phe Ala Glu Arg Phe 275 280 285

Ala Asp Ala Tyr Lys Asn Glu Cys Ile Ala Phe Val Glu Leu Ile Leu

290 295 300 Gly Glu Arg Glu Asn Pro Cys Thr Pro Ala Asp Ala Val Ala Ala Ala 305 310 315 Ile Val Ala Asp Ala Ala Gln Leu Ser Leu Val Thr Gly Glu Pro Val 325 330 Lys Ile Pro Thr Val Arg Glu Ile Leu Glu Gly Ser Ala Gln Pro Val Glu Val Arg Ala Leu Val Pro Ser Ala 355 <210> 303 <211> 1146 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1123) <223> RXN01406 <400> 303 gttcctcatt cctctaatcg gcgcactatc tttgcctcgc gacggcggtg cccgagcctt 60 ttcctcctct tagaaaccca cttctgaaag gtataaaaac atg act att cga atc Met Thr Ile Arg Ile gga ctc gtt ggc tac ggt gtc ggc ggc agg ctc ttt cac acc cct tac Gly Leu Val Gly Tyr Gly Val Gly Gly Arg Leu Phe His Thr Pro Tyr 10 atc caa gct tct acg cac tgc gaa tta gta ggc gta gtt gct cgt tcc 211 Ile Gln Ala Ser Thr His Cys Glu Leu Val Gly Val Val Ala Arg Ser gaa ggc acc aaa gca gcc gtt gca gaa gat ctt cca gat gtt gcc atc 259 Glu Gly Thr Lys Ala Ala Val Ala Glu Asp Leu Pro Asp Val Ala Ile 45 gtg gga tcg ctg aca gaa ctc ctc gaa ctg ggc gtc gat gca gtg gtg 307 Val Gly Ser Leu Thr Glu Leu Leu Glu Leu Gly Val Asp Ala Val Val atc tee acc eet eea gee acg ege egg gaa etg gee ttg gaa gea ate Ile Ser Thr Pro Pro Ala Thr Arg Arg Glu Leu Ala Leu Glu Ala Ile 80 aac gca ggt gtc gca gtg gtt gcc gat aaa ccg ttt gca cca tca gcc 403 Asn Ala Gly Val Ala Val Val Ala Asp Lys Pro Phe Ala Pro Ser Ala

gca gat gcc atg gaa ctt gtc gaa gcc gcc gaa aag gct gga gtg ctg

Ala Asp Ala Met Glu Leu Val Glu Ala Ala Glu Lys Ala Gly Val Leu 105 110 115

95

451

PCT/IB00/00943 WO 01/00844

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cta gac ctg a Leu Asp Leu 1 150				Gly Pro Glu		595
ttg ctg cgc (Leu Leu Arg)						643
atg ggg ccg (Met Gly Pro A		Val Thr A			Asp Leu	691
cca gaa ggc o Pro Glu Gly 1 200						739
tcg ggc gcc g Ser Gly Ala v 215						787
tcc tgg gaa a Ser Trp Glu 1 230				Ser Tyr Val		835
tac acc gac of Tyr Thr Asp						883
aat gac cgc (Asn Asp Arg (Gly Tyr G			Gly Thr	931
ttg gtt acc g Leu Val Thr 2 280						979
tac acc cgc t Tyr Thr Arg 1 295						1027
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Pro Asp Val Ala Ile Val Gly Ser Leu Thr Glu Leu Leu Glu Leu Gly 50 55 60

Val Asp Ala Val Val Ile Ser Thr Pro Pro Ala Thr Arg Arg Glu Leu 65 70 75 80

Ala Leu Glu Ala Ile Asn Ala Gly Val Ala Val Val Ala Asp Lys Pro 85 90 95

Phe Ala Pro Ser Ala Ala Asp Ala Met Glu Leu Val Glu Ala Ala Glu 100 105 110

Lys Ala Gly Val Leu Leu Asn Val Phe His Asn Arg Arg Asn Asp Thr 115 120 125

His Ile Val Thr Ala Leu Gly Ile Gln Glu Glu Leu Gly Ala Met Arg 130 135 140

Gly Leu Asp Leu Arg Leu Asp Leu Ile Glu Pro Asp Ser Leu Glu Ala 145 150 155 160

Gly Pro Glu Gly Gly Leu Leu Arg Asp Leu Gly Ser His Val Val Asp 165 170 175

Gln Thr Leu Val Leu Met Gly Pro Ala Thr Ser Val Thr Ala Gln Leu 180 185 190

Gly Ser Ile Asp Leu Pro Glu Gly Pro Thr Asn Ala Arg Phe Arg Ile 195 200 205

Val Leu Glu His Glu Ser Gly Ala Val Ser His Ile Ser Ala Ser Lys 210 215 220

Ile Asp Arg Leu Glu Ser Trp Glu Ile Arg Leu Val Gly Glu Arg Gly 225 230 235 240

Ser Tyr Val Ser Asn Tyr Thr Asp Val Gln Thr Val Ala Ile Lys Gln 245 250 255

Gly Leu Arg Pro Thr Asn Asp Arg Glu His Trp Gly Tyr Glu Ser Glu 260 265 270

Glu Arg Trp Gly Thr Leu Val Thr Asp Glu Gly Ser Lys Val Ile Pro 275 280 285

Ser Ala Gln Gly Asp Tyr Thr Arg Phe Tyr Asp Ala Phe Ala Leu Ala 290 295 300

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140

135

agc Ser 150	Val	caa Gln	gca Ala	cgc Arg	ggc Gly 155	gcc Ala	gca Ala	aaa Lys	gta Val	ggt Gly 160	gag Glu	cat His	atc	acc Thr	gag Glu 165	595
cac His	ctc Leu	aac Asn	caa Gln	ccc Pro 170	gca Ala	gac Asp	atg Met	ggc Gly	ggt Gly 175	gtg Val	ttg Leu	tgg Trp	att Ile	ctt Leu 180	ggc Gly	643
tgc Cys	cac His	atg Met	ctc Leu 185	gat Asp	gca Ala	ttg Leu	gtg Val	gaa Glu 190	gtc Val	ttc Phe	gga Gly	gct Ala	cca Pro 195	gaa Glu	tcc Ser	691
gtg Val	aac Asn	gcc Ala 200	cga Arg	gtg Val	cat His	aag Lys	acc Thr 205	gca Ala	aaa Lys	ctc Leu	tct Ser	gac Asp 210	gac Asp	acc Thr	agc Ser	739
cgc Arg	gaa Glu 215	gac Asp	tca Ser	gcc Ala	tcc Ser	gca Ala 220	ctg Leu	ctg Leu	tac Tyr	tac Tyr	cca Pro 225	gat Asp	ttc Phe	tcc Ser	gtc Val	787
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cga Arg	ctc Leu	acg Thr	gtc Val	tat Tyr 250	gga Gly	acc Thr	aag Lys	ggc Gly	atg Met 255	atc Ile	gaa Glu	gcc Ala	gga Gly	atc Ile 260	ctc Leu	883
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			cgc Arg													1075
gga Gly	tcc Ser	cgc Arg	aat Asn	gtg Val 330	gcg Ala	cca Pro	gtt Val	gag Glu	gat Asp 335	gct Ala	ctc Leu	aca Thr	gtc Val	gct Ala 340	cgc Arg	1123
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Val Val Ala Ala Asp Thr Asp Ser Arg Leu Gln Tyr Phe Thr Asp 35 40 45

Lys Tyr Asp Val Glu Pro Arg Glu Ile Asp Asp Val Leu Asn Asp Asp 50 55 60

Arg Ile Asn Ala Ile Met Val His Ser Lys Ser Lys Asp Met Val Pro 65 70 75 80

His Ala Lys Arg Ala Leu Ala Ala Gly Lys Ser Val Val Val Glu Lys 85 90 95

Pro Gly Gly Gly Thr Val Ala Asp Leu Glu Glu Leu Leu Ala Leu Lys 100 105 110

Glu Ala Ala Asp Pro Gln Arg Ile Val Gln Val Gly Tyr Asn Val Arg 115 120 125

Leu Ser Glu Ser Val Gln Arg Leu Lys Glu Leu Leu Asp Ala Gly Leu 130 135 140

Ile Gly Glu Val Val Ser Val Gln Ala Arg Gly Ala Ala Lys Val Gly 145 150 155 . 160

Glu His Ile Thr Glu His Leu Asn Gln Pro Ala Asp Met Gly Gly Val 165 170 175

Leu Trp Ile Leu Gly Cys His Met Leu Asp Ala Leu Val Glu Val Phe 180 185 190

Gly Ala Pro Glu Ser Val Asn Ala Arg Val His Lys Thr Ala Lys Leu 195 200 205

Ser Asp Asp Thr Ser Arg Glu Asp Ser Ala Ser Ala Leu Leu Tyr Tyr 210 215 220

Pro Asp Phe Ser Val Ser Phe Ser Phe Asp Gly His Asp Asp Leu Glu 225 230 235 240

Trp Phe Glu Ser Ser Arg Leu Thr Val Tyr Gly Thr Lys Gly Met Ile 245 250 255

Glu Ala Gly Ile Leu Pro Gln Thr Leu Arg Val Tyr Leu Asn Glu Ser 260 265 270

Arg Gln Gly Trp Pro Gln Gly Trp Thr Glu Trp Thr Gln Ser Tyr Phe 275 280 285

Thr Pro Pro Phe Ala Arg Thr Glu Ser Asn Lys Phe Ser Glu Leu Pro 290 295 300

Glu Leu Glu Asn Ile Ser Asn Phe Arg Thr Glu Met Gln Gly Trp Val

Ası	n Ser	: Ile	e Arç	325		/ Ser	Arg	Asr	Val 330		a Pro	Val	l Gli	335	Ala	
Let	Thr	Val	Ala 340		ı Ile	· Val	. Ser	Ala 345		з Туг	Glu	Sei	350		n Asn	
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cag Gln	ggt Gly	gtg Val	gaa Glu 25	tat Tyr	tac Tyr	cga Arg	aat Asn	gcg Ala 30	gat Asp	cct Pro	tcc Ser	gaa Glu	act Thr 35	Val	ccg Pro	211
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gcc Ala 70	gac Asp	gcc Ala	acc Thr	gag Glu	gct Ala 75	tca Ser	caa Gln	aac Asn	tgc Cys	act Thr 80	atc Ile	aaa Lys	atc Ile	gcc Ala	gat Asp 85	355
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ctg Leu	ggc Gly	gat Asp	cat His 105	tac Tyr	cgc Arg	gcg Ala	acc Thr	atc Ile 110	gac Asp	gag Glu	tcc Ser	acc Thr	gcc Ala 115	gag Glu	cca Pro	451
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						ctt Leu										739
						gtc Val 220										787
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cgc Arg	aac Asn	gtc Val	cac His 265	atc Ile	gga Gly	cca Pro	tcc Ser	gac Asp 270	cac His	gtc Val	caa Gln	tgg Trp	ctc Leu 275	gat Asp	gac Asp	931
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Ser Glu Thr Val Pro Gly Leu Met His Val Lys Phe Gly Asp Tyr His
35 40 45

Val Gly Asp Ile Glu Phe Val Ala Ala Phe Asp Val Asp Ala Glu Lys 50 55 60

Val Gly Ile Asp Leu Ala Asp Ala Thr Glu Ala Ser Gln Asn Cys Thr 65 70 75 80

Ile Lys Ile Ala Asp Val Pro Gln Thr Gly Ile Asn Val Leu Arg Gly
85 90 95

Pro Thr Leu Asp Gly Leu Gly Asp His Tyr Arg Ala Thr Ile Asp Glu 100 105 110

Ser Thr Ala Glu Pro Val Asp Val Val Gln Ala Leu Ile Asp Ala Lys 115 120 125

Ala Asp Val Leu Val Ser Tyr Leu Pro Val Gly Ser Glu Glu Ala Asp 130 135 140

Lys Phe Tyr Ala Gln Ala Ala Ile Asp Ala Gly Cys Ala Phe Val Asn 145 150 155 160

Ala Leu Pro Val Phe Ile Ala Ser Asp Pro Glu Trp Ala Lys Lys Phe 165 170 175

Thr Asp Ala Gly Ile Pro Ile Val Gly Asp Asp Ile Lys Ser Gln Ile 180 185 190

Gly Ala Thr Ile Thr His Arg Val Leu Ala Arg Leu Phe Glu Glu Arg 195 200 205

Gly Val Arg Val Asp Arg Thr Met Gln Leu Asn Val Gly Gly Asn Met 210 225 220

Asp Phe Lys Asn Met Leu Asp Arg Asn Arg Leu Glu Ser Lys Lys Val 225 230 235 240

Ser Lys Thr Gln Ala Val Thr Ser Asn Ile Pro Asp Gly Pro Leu Ser 245 250 255

Gly Lys Val Glu Asp Arg Asn Val His Ile Gly Pro Ser Asp His Val 260 265 270

Gin Trp Leu Asp Asp Arg Lys Trp Ala Tyr Val Arg Leu Glu Gly Thr 275 280 285

Ala Phe Gly Gly Val Pro Leu Asn Leu Glu Tyr Lys Leu Glu Val Trp

295 290 300 Asp Ser Pro Asn Ser Ala Gly Ile Ile Ile Asp Ala Val Arg Ala Ala 310 315 Lys Ile Ala Leu Asp Arg Gly Ile Gly Gly Pro Ile Met Pro Ala Ser 325 Ser Tyr Leu Met Lys Ser Pro Pro Glu Gln Leu Pro Asp Asp Val Ala 340 345 Cys Glu Arg Leu Glu Ala Phe Ile Ile Glu Ala <210> 309 <211> 795 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(772) <223> RXN03057 <400> 309 catcaacqcc qaqtacaact aaqqacaact qataatqaca aatqctqcaa ttqtcqqatq 60 aggagacgtc gcaaccgttc atacagaagc gctggaagct ttg gct tcc gat ctt Leu Ala Ser Asp Leu ggt att aag ttc gtc gca gtg gtg gat aaa gat cta gag act gct gag 163 Gly Ile Lys Phe Val Ala Val Val Asp Lys Asp Leu Glu Thr Ala Glu 10 aaa ttt geg aeg gga ett gga get get gge gat tet tea gaa age age 211 Lys Phe Ala Thr Gly Leu Gly Ala Ala Gly Asp Ser Ser Glu Ser Ser 25 gtc aag gcc cac ggc agc ctg ccg gct ttg ttc tcc aaa aag aag atc 259 Val Lys Ala His Gly Ser Leu Pro Ala Leu Phe Ser Lys Lys Ile 40 45 gat gtt cta cac atc acc ccc cac gac caa cac att ggt ttg gct 307 Asp Val Leu His Ile Thr Thr Pro His Asp Gln His Ile Gly Leu Ala 60 ctc gaa gcg cta cac cac ggt gta aat gtc atc ctg gaa aag ccg ttg Leu Glu Ala Leu His His Gly Val Asn Val Ile Leu Glu Lys Pro Leu gct aat gag ttg gac cag gcg cag cgt ctc atc gac tac ttg gat gaa 403 Ala Asn Glu Leu Asp Gln Ala Gln Arg Leu Ile Asp Tyr Leu Asp Glu aac ccc gat ggt cca aag att gca gtg tgc tat cag aac cgt tac aac Asn Pro Asp Gly Pro Lys Ile Ala Val Cys Tyr Gln Asn Arg Tyr Asn 105 110

gtt tee tee cag gaa etg egt egt etg ete gat tea ggt gae ete ggt

Val Ser Ser Gln Glu Leu Arg Arg Leu Leu Asp Ser Gly Asp Leu Gly 125 gcc atc aat ggt gca tat tcc tct gtg gtg tgg acc cgc acc cca ggc Ala Ile Asn Gly Ala Tyr Ser Ser Val Val Trp Thr Arg Thr Pro Gly tac tac acc cag aaa cct tgg cgt ggc cag caa gca cac tcc ggt ggt 595 Tyr Tyr Thr Gln Lys Pro Trp Arg Gly Gln Gln Ala His Ser Gly Gly 155 ggc ctg ctg atg aac caa gca att cac acc ctg gat ctg ctg cag tgg 643 Gly Leu Leu Met Asn Gln Ala Ile His Thr Leu Asp Leu Leu Gln Trp 170 175 ttc ctt gga aag gca aca gaa gtc aag ggc act gtc tcc acc gat aag 691 Phe Leu Gly Lys Ala Thr Glu Val Lys Gly Thr Val Ser Thr Asp Lys 190 tat gcc gat gtc atc gat gtt gaa gac acc gca cac gca tac atc ggt 739 Tyr Ala Asp Val Ile Asp Val Glu Asp Thr Ala His Ala Tyr Ile Gly 205 cac gag tcc gga gtc cac acc agt gaa gtg agt tgaaccatgc tattggtgat 792 His Glu Ser Gly Val His Thr Ser Glu Val Ser 215 220 aca 795 <210> 310 <211> 224 <212> PRT <213> Corynebacterium glutamicum <400> 310 Leu Ala Ser Asp Leu Gly Ile Lys Phe Val Ala Val Val Asp Lys Asp 10 Leu Glu Thr Ala Glu Lys Phe Ala Thr Gly Leu Gly Ala Ala Gly Asp 25 Ser Ser Glu Ser Ser Val Lys Ala His Gly Ser Leu Pro Ala Leu Phe 35 Ser Lys Lys Ile Asp Val Leu His Ile Thr Thr Pro His Asp Gln His Ile Gly Leu Ala Leu Glu Ala Leu His His Gly Val Asn Val Ile Leu Glu Lys Pro Leu Ala Asn Glu Leu Asp Gln Ala Gln Arg Leu Ile Asp Tyr Leu Asp Glu Asn Pro Asp Gly Pro Lys Ile Ala Val Cys Tyr 105 110 Gln Asn Arg Tyr Asn Val Ser Ser Gln Glu Leu Arg Arg Leu Leu Asp 120 Ser Gly Asp Leu Gly Ala Ile Asn Gly Ala Tyr Ser Ser Val Val Trp

130 135 140 Thr Arg Thr Pro Gly Tyr Tyr Thr Gln Lys Pro Trp Arg Gly Gln Gln 150 Ala His Ser Gly Gly Gly Leu Leu Met Asn Gln Ala Ile His Thr Leu 170 Asp Leu Leu Gln Trp Phe Leu Gly Lys Ala Thr Glu Val Lys Gly Thr Val Ser Thr Asp Lys Tyr Ala Asp Val Ile Asp Val Glu Asp Thr Ala His Ala Tyr Ile Gly His Glu Ser Gly Val His Thr Ser Glu Val Ser 215 <210> 311 <211> 795 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(772) <223> FRXA02902 <400> 311 catcaacgcc gagtacaact aaggacaact gataatgaca aatgctgcaa ttgtcggatg 60 aggagacgtc gcaaccgttc atacagaagc gctggaagct ttg gct tcc gat ctt Leu Ala Ser Asp Leu 1 ggt att aag ttc gtc gca gtg gtg gat aaa gat cta gag act gct gag 163 Gly Ile Lys Phe Val Ala Val Val Asp Lys Asp Leu Glu Thr Ala Glu 211 aaa ttt gcg acg gga ctt gga gct gct ggc gat tct tca gaa agc agc Lys Phe Ala Thr Gly Leu Gly Ala Ala Gly Asp Ser Ser Glu Ser Ser gtc aag gcc cac ggc agc ctg ccg gct ttg ttc tcc aaa aag aag atc Val Lys Ala His Gly Ser Leu Pro Ala Leu Phe Ser Lys Lys Ile 45 307 gat gtt cta cac atc acc ccc cac gac caa cac att ggt ttg gct Asp Val Leu His Ile Thr Thr Pro His Asp Gln His Ile Gly Leu Ala 355 ctc gaa gcg cta cac cac ggt gta aat gtc atc ctg gaa aag ccg ttg Leu Glu Ala Leu His His Gly Val Asn Val Ile Leu Glu Lys Pro Leu gct aat gag ttg gac cag gcg cag cgt ctc atc gac tac ttg gat gaa Ala Asn Glu Leu Asp Gln Ala Gln Arg Leu Ile Asp Tyr Leu Asp Glu

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90

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gtt Val	tcc Ser	tcc Ser 120	cag Gln	gaa Glu	ctg Leu	cgt Arg	cgt Arg 125	ctg Leu	ctc Leu	gat Asp	tca Ser	ggt Gly 130	gac Asp	ctc Leu	ggt Gly	499
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tac Tyr 150	tac Tyr	acc Thr	cag Gln	aaa Lys	cct Pro 155	tgg Trp	cgt Arg	ggc Gly	cag Gln	caa Gln 160	gca Ala	cac His	tcc Ser	ggt Gly	ggt Gly 165	595
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tat Tyr	gcc Ala	gat Asp 200	gtc Val	atc Ile	gat Asp	gtt Val	gaa Glu 205	gac Asp	acc Thr	gca Ala	cac His	gca Ala 210	tac Tyr	atc Ile	ggt Gly	739
cac His	gag Glu 215	tcc Ser	gga Gly	gtc Val	cac His	acc Thr 220	agt Ser	gaa Glu	gtg Val	agt Ser	tgaa	accat	igc 1	tatto	ggtgat	792
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gat Asp	cto Lev	aac Asn	gto Val 105	. Ile	gtc Val	ccg Pro	gcc Ala	gag Glu 110	Let	g agt 1 Ser	cgc Arg	caa Gln	cto Lev	ttg Leu	ccc Pro	451
gcc Ala	Leu	Arg 120	Ala	gca Ala	tcc Ser	ggc Gly	tgc Cys 125	Val	ato Ile	tac Tyr	atc Ile	aac Asn 130	Ser	ggc Gly	gcc Ala	499
GTÀ	135	GIY	Pro	His	Pro	Gly 140	Asn	Thr	Ile	Tyr	Ala 145	Ala	Ser	aaa Lys	His	547
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Ala	215	Glu	Thr	Thr	Gln	11e 220	Thr	Asn	Val	Asp	Val 225	Arg	Pro	cgt Arg	atc Ile	787
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Glu	Pro 50	Ile	Glu	Ser	Asp	Ile 55	Val	Lys	Glu	Val	Leu (Glu (Glu	Gly	Gly	

Val Asp Lys Leu Lys Asn Leu Asp His Val Asp Thr Leu Val His Ala Ala Ala Val Ala Arg Asp Thr Thr Ile Glu Ala Gly Ser Val Ala Glu 90 Trp His Ala His Leu Asp Leu Asn Val Ile Val Pro Ala Glu Leu Ser 100 105 Arg Gln Leu Leu Pro Ala Leu Arg Ala Ala Ser Gly Cys Val Ile Tyr 120 Ile Asn Ser Gly Ala Gly Asn Gly Pro His Pro Gly Asn Thr Ile Tyr 130 Ala Ala Ser Lys His Ala Leu Arg Gly Leu Ala Asp Ala Phe Arg Lys Glu Glu Ala Asn Asn Gly Ile Arg Val Ser Thr Val Ser Pro Gly Pro 165 Thr Asn Thr Pro Met Leu Gln Gly Leu Met Asp Ser Gln Gly Thr Asn 180 185 Phe Arg Pro Glu Ile Tyr Ile Glu Pro Lys Glu Ile Ala Asn Ala Ile 205 Arg Phe Val Ile Asp Ala Gly Glu Thr Thr Gln Ile Thr Asn Val Asp 215 Val Arg Pro Arg Ile Glu Leu Ala Asp Arg Lys Asp 230 <210> 315 <211> 1008 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(985) <223> RXN02654 <400> 315 tattttcgga aatttataca gcaatcctcg aaatcctaat aaagatccct tatcgtggga 60 gaggtacggt agttcgttcg aggacaacgt cgagaaaggc atg att tca ttg cta Met Ile Ser Leu Leu aat gat cca cgt acg cta ttc ccg aaa gtc gat ccc cca aag caa agc Asn Asp Pro Arg Thr Leu Phe Pro Lys Val Asp Pro Pro Lys Gln Ser 10 caq ccg gaa cca ggc cta gat ata aaa ctt tcc ccc caa gcc gat att Gln Pro Glu Pro Gly Leu Asp Ile Lys Leu Ser Pro Gln Ala Asp Ile 25 ggt ctc tcc agc tat caa gga agt gga agg ctt aag ggc cgc aag gct

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tat Tyr 70	Ala	cgc Arg	gag Glu	Gly ggg	gca Ala 75	Asp	gtt Val	gcg Ala	atc	gct Ala 80	Tyr	ttg Leu	Pro	gaa Glu	gaa Glu 85	355
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cac His	cct Pro	Val	gag Glu 265	ttg Leu	gca Ala	ggt Gly	gcg Ala	tac Tyr 270	gtt Val	ttt Phe	ctc Leu	Ala	tct Ser 275	gac Asp	gaa Glu	931
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Lys Gly Arg Lys Ala Leu Ile Thr Gly Gly Asp Ser Gly Ile Gly Ala 50 55 60

Ala Val Ala Ile Ala Tyr Ala Arg Glu Gly Ala Asp Val Ala Ile Ala 65 70 75 80

Tyr Leu Pro Glu Glu Gln Ala Asp Ala Asp Arg Val Leu Gln Ala Ile 85 90 95

Glu Glu Thr Gly Gln Lys Ala Phe Ser Phe Pro Gly Asp Leu Arg Asp 100 105 110

Pro Glu Tyr Cys Arg Ser Leu Val Gln Glu Thr Val Asn Ala Leu Gly
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Gly Leu Asp Ile Leu Val Asn Asn Ala Ser Arg Gln Val Trp Ala Pro 130 135 140

Gly Leu Thr Glu Ile Thr Asp Glu Asn Phe Asp Gln Thr Leu Gln Val 145 150 155 160

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Lys Pro Gly Ser Ser Ile Ile Phe Thr Ser Ser Ile Gln Ala Tyr Gln 180 185 190

Pro Ser Glu Thr Leu Leu Asp Tyr Ala Met Thr Lys Ala Ala Leu Asn 195 200 205

Asn Leu Ser Lys Gly Leu Ala Ser Ser Leu Ile Gly Asp Gly Ile Arg 210 215 220

Val Asn Ser Val Ala Pro Gly Pro Phe Trp Thr Pro Leu Gln Pro Ser 225 230 235 240

His Gly Gln Pro Gln Glu Lys Ile Glu Gly Phe Gly Gln His Ala Pro 245 250 255

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Lys Gly Arg Lys Ala Leu Ile Thr Gly Gly Asp Ser Gly Ile Gly Ala

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- Glu Glu Thr Gly Gln Lys Ala Phe Ser Phe Pro Gly Asp Leu Arg Asp 100 105 110
- Pro Glu Tyr Cys Arg Ser Leu Val Gln Glu Thr Val Asn Ala Leu Gly
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- Gly Leu Asp Ile Leu Val Asn Asn Ala Ser Arg Gln Val Trp Ala Pro 130 135 140
- Gly Leu Thr Glu Ile Thr Asp Glu Asn Phe Asp Gln Thr Leu Gln Val 145 150 155 160
- Asn Leu Tyr Gly Ser Phe Arg Val Thr Lys Ala Ala Ile Pro His Leu 165 170 175
- Lys Pro Gly Ser Ser Ile Ile Phe Thr Ser Ser Ile Gln Ala Tyr Gln 180 185 190
- Pro Ser Glu Thr Leu Leu Asp Tyr Ala Met Thr Lys Ala Ala Leu Asn 195 200 205
- Asn Leu Ser Lys Gly Leu Ala Ser Ser Leu Ile Gly Asp Gly Ile Arg 210 215 220
- Val Asn Ser Val Ala Pro Gly Pro Phe Trp Thr Pro Leu Gln Pro Ser 225 230 235 240
- His Gly Gln Pro Gln Glu Lys Ile Glu Gly Phe Gly Gln His Ala Pro 245 250 255
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Gln Val Val Ser Glu Ile Thr Ser Val Ile Asn Gly Ile Leu Asn Ala 65 70 75 80

Ala Asp His His Asn Ile Lys Asp Gln Ile Ala Ala Val Ala Leu Asp 85 90 95

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Thr Pro Cys Ile Thr Tyr Ala Asp Ser Arg Ser Ala Gln Tyr Val Glu 115 120 125

Gln Leu Arg Ala Glu Ile Asp Glu Lys Ala Tyr His Gly Arg Thr Gly 130 135 140

Val Cys Leu His Thr Ser Tyr His Pro Ser Arg Leu Leu Trp Leu Lys 145 150 155 160

Thr Glu Phe Glu Lys Glu Phe Asn Lys Ala Lys Tyr Val Met Thr Ile 165 170 175

Gly Glu Tyr Val Tyr Phe Lys Leu Ala Gly Ile Thr Gly Met Ala Thr 180 185 190

Ser Ile Ala Ala Trp Ser Gly Ile Leu Asp Ala His Thr Gly Glu Leu 195 200 205

Asp Leu Thr Ile Leu Glu His Ile Gly Val Asp Pro Ala Leu Phe Gly 210 215 220

Glu Ile Arg Asn Pro Asp Glu Pro Ala Thr Asp Ala Lys Val Val Asp 225 230 235 240

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- Phe Ser Gly Glu Arg Ser Ile Gly Trp Ala Ala Ser Ala Gln Ala Thr 355 360 365
- Ile Thr Asn Ile Gln Glu Gln Thr Gly Pro Glu His Leu Trp Arg Gly 370 380
- Val Phe Glu Ala Leu Ala Leu Ser Tyr Gln Arg Val Trp Glu His Met 385 390 395 400
- Gly Lys Ala Gly Ala Ala Pro Glu Arg Val Ile Ala Ser Gly Arg Val 405 410 415
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- Thr Pro Val Ile Pro Leu Glu Met Lys Arg Ala Thr Leu Arg Gly Thr 435 440 445
- Ala Leu Ile Val Leu Glu Gln Leu Glu Pro Gly Gly Thr Arg Ala Thr 450 455 460
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Gly Ala Gln Ala Ala Ala Leu Ala Glu Ala Ile Gly Gly Glu Gly 145 150 155 160

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Arg Gly Gln Gly Phe Glu Glu Glu Ile Ala Lys His Glu Gly Ile Ser 180 185 190

Ile Val Ala Lys Gln Thr Ala Asn Phe Asp Arg Gly Glu Gly Leu Asp 195 200 205

Val Ala Thr Asn Leu Leu Gln Ala His Pro Asn Val Lys Ala Ile Phe 210 215 220

Ala Glu Asn Asp Glu Met Ala Leu Gly Ala Ile Glu Ala Leu Gly Ala 225 230 235 240

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Phe Trp Glu Ser Pro Glu Glu Ala Thr Lys Gln Ala Glu Trp Ala Leu 180 185 190

Gln His Ser Thr Val Ala Val Gly Asn Lys Glu Glu Cys Glu Ile Ala 195 200 205

Val Gly Glu Thr Glu Pro Glu Arg Ala Gly Arg Ala Leu Leu Glu Arg 210 215 220

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Met Thr Lys Asp Glu Thr Val Glu Val Pro Pro Phe Phe Val Asp Val 245 250 255

Ile Asn Gly Leu Gly Ala Gly Asp Ala Phe Gly Gly Ala Leu Cys His 260 265 270

Gly Leu Leu Ser Glu Trp Pro Leu Glu Lys Val Leu Arg Phe Ala Asn 275 280 285

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Cys Glu Pro Leu Leu Asn Pro Val Asp Met Trp Arg Glu Asp Asn Pro

90 95 100

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Gly Ser His Tyr Pro Gly Tyr Ser Pro Leu Ala Pro Glu Ser Glu Gly 65 70 75 80

Lys Asp Ala Glu Lys Cys Glu Pro Leu Leu Asn Pro Val Asp Met Trp 85 90 95

Arg Glu Asp Asn Pro Ile Thr Gly Val Pro Phe Thr Glu Pro Val Leu 100 105 110

Ala Thr Ser Ser Thr Glu Asn Ala Ile Asn Leu Arg Asn Gln Arg Tyr 115 120 125

Leu Ile Val Arg Asp Asn Leu Pro Ala Arg Gly Leu Ala Thr Trp Thr 130 135 140

Ala Phe Ala Ser Asn Pro Arg Asn His Val Ala Leu Val Ala Gln Phe 165 170 175

Gly Val Asn Glu Ser Ala Gly Val Phe Ser Glu Trp Pro Gly Glu Leu 180 185 190

Gly Leu Ala Ala Leu Arg Asp Ala Glu Leu Met Glu Thr Phe Gly Thr 195 200 205

Glu Ala Ala Lys Glu Trp Arg Ala Gly Gly Val His Lys Leu Tyr Gly 210 215 220

Tyr Met Ala Asp Leu Ala Ser Glu Pro Arg Trp Ser Arg Phe Asn Gly
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Val Arg Gly Leu Gln Gly Pro Glu Leu Ser Lys Asn Ser Val Ser Thr 260 265 270

Thr Ile Lys His Phe Pro Gly Gly Gly Val Arg Leu Asp Gly His Asp 275 280 285

Pro His Phe His Trp Gly Gln Thr Asn Glu Tyr Pro Thr Glu Asp Ala 290 295 300

Leu Gly Lys Tyr His Leu Pro Pro Phe Gln Ala Ala Ile Asp Ala Gly 305 310 315 320

Cys Ala Ser Ile Met Pro Tyr Tyr Ala Arg Pro Met Asn Asn Ser Ala 325 330 335

Asn Gln Leu Asp Gln Gln Leu Trp Gln Asn Pro Thr Thr Gln Phe Glu 340 345 350

Glu Val Ala Phe Ala Tyr Asn Arg Thr Phe Ile Gln Asp Leu Leu Arg 355 360 365

Asp Ala Met Gly His Arg Gly Tyr Val Asn Ser Asp Ser Gly Val Ile 370 375 380

Asp Ala Met Met Trp Gly Val Glu Glu Leu Ser Glu Pro Glu Arg Phe 385 390 395 400

Ala Ala Val Arg Ala Gly Thr Asp Ile Phe Ser Asp Met Ala Asn 405 410 415

Pro Arg Arg Leu Leu Glu Ala Val Ala Glu Gly His Leu Asp Glu Ser 420 425 430

Glu Leu Asn Gln Pro Val Gln Arg Leu Leu Glu Glu Ile Phe Gln Leu 435 440 445

Gly Leu Phe Glu Asn Pro Tyr Val Ser Glu Asp Glu Ala Glu Lys Ile 450 455 460

Ile Gly Ala Pro Glu Val Ser Ala Leu Gly Asn Lys Ala Gln Leu Asp 465 470 475 480

Ser Val Thr Leu Leu Arg Asn Asn Pro Ile Arg Ala Ala Thr Gly Ser 485 490 495

Cys Ser Lys Pro Glu Asp Leu Pro Ile Gly Tyr Trp Pro Tyr Gln Asp
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Pro Gly Tyr Ser Pro Leu Ala Pro Glu Ser Glu Gly Lys Asp Ala Glu 50 55 60

Lys Cys Glu Pro Leu Leu Asn Pro Val Asp Met Trp Arg Glu Asp Asn 65 70 75 80

Pro Ile Thr Gly Val Pro Phe Thr Glu Pro Val Leu Ala Thr Ser Ser 85 90 95

Thr Glu Asn Ala Ile Asn Leu Arg Asn Gln Arg Tyr Leu Ile Val Arg 100 105 110

Asp Asn Leu Pro Ala Arg Gly Leu Ala Thr Trp Thr Asn Ala Val Gln 115 120 125

Glu Val Ala Glu Arg Ser Arg Leu Gly Ile Pro Val Ala Phe Ala Ser 130 135 140

Asn Pro Arg Asn His Val Ala Leu Val Ala Gln Phe Gly Val Asn Glu 145 150 155 160

Ser Ala Gly Val Phe Ser Glu Trp Pro Gly Glu Leu Gly Leu Ala Ala 165 170 175

Leu Arg Asp Ala Glu Leu Met Glu Thr Phe Gly Thr Glu Ala Ala Lys 180 185 190

Glu Trp Arg Ala Gly Gly Val His Lys Leu Tyr Gly Tyr Met Ala Asp 195 200 205

Leu Ala Ser Glu Pro Arg Trp Ser Arg Phe Asn Gly Thr Phe Gly Glu 210 215 220

Asp Pro Glu Leu Ile Ser Asp Tyr Ile Ala Ala Val Val Arg Gly Leu 225 230 235 240

Gln Gly Pro Glu Leu Ser Lys Asn Ser Val Ser Thr Thr Ile Lys His

245 250 255

Phe Pro Gly Gly Val Arg Leu Asp Gly His Asp Pro His Phe His 260 265 270

Trp Gly Gln Thr Asn Glu Tyr Pro Thr Glu Asp Ala Leu Gly Lys Tyr 275 280 285

His Leu Pro Pro Phe Gln Ala Ala Ile Asp Ala Gly Cys Ala Ser Ile 290 295 300

Met Pro Tyr Tyr Ala Arg Pro Met Asn Asn Ser Ala Asn Gln Leu Asp 305 310 315 320

Gln Gln Leu Trp Gln Asn Pro Thr Thr Gln Phe Glu Glu Val Ala Phe 325 330 335

Ala Tyr Asn Arg Thr Phe Ile Gln Asp Leu Leu Arg Asp Ala Met Gly 340 345 350

His Arg Gly Tyr Val Asn Ser Asp Ser Gly Val Ile Asp Ala Met Met 355 360 365

Trp Gly Val Glu Glu Leu Ser Glu Pro Glu Arg Phe Ala Ala Val 370 375 380

Arg Ala Gly Thr Asp Ile Phe Ser Asp Met Ala Asn Pro Arg Arg Leu 385 390 395 400

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405 410 415

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Asn Pro Tyr Val Ser Glu Asp Glu Ala Glu Lys Ile Ile Gly Ala Pro 435 440 445

Glu Val Ser Ala Leu Gly Asn Lys Ala Gln Leu Asp Ser Val Thr Leu 450 455 460

Leu Arg Asn Asn Pro Ile Arg Ala Ala Thr Gly Ser Cys Ser Lys Pro 465 470 475 480

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170

190

Phe Gly Leu Ile Thr Ala Ala Leu Ile Ser Arg Lys Glu Ser Gly Ser

185

165

180

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Ser Gly Asn Arg Thr Asp Ile Leu Ser Phe Ile Arg Asp Arg Gly Ile 450 455 460

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Val Ala Glu Ala Asn Ser Val Gln Ala Gln Asp Val Val Met Glu Ser 100 105 110

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Phe Gln Leu Thr Glu Ser Val Leu Gly Gly Ser Gly Met Asn Ile Ala 165 170 175

Ala Leu Val Gly Glu Glu Ser Leu Ser Thr Thr Gln Glu Arg Met Arg 180 185 190

Gly Ile Ser His Ala Ala Ser Ile Tyr Gly Ala Glu Val Thr Phe His 195 200 205

Phe Gly His Tyr Ser Val Glu Ser Gly Glu Met Ala Gln Val Val 210 215 220

Phe Asn Asn Gly Leu Pro Asp Ala Leu Ile Val Ala Ser Pro Arg Leu 225 230 235 240

Met Ala Gly Val Met Arg Ala Phe Thr Arg Leu Asn Val Arg Val Pro 245 250 255

His Asp Val Val Ile Gly Gly Tyr Asp Asp Pro Glu Trp Tyr Ser Phe 260 265 270

Val Gly Ala Gly Ile Thr Thr Phe Val Pro Pro His Glu Glu Met Gly 275 280 285

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ctg cgc gtc atc Leu Arg Val Ile					
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Glu Pro Val Glu 50	Val Phe Gly 55	Tyr Thr Asn	Ser Phe Lys 60	His Gly	Asp

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Gly Ala Leu Ala Thr Ile Ser Ala Thr Thr Ala Ala Glu Pro Ala Leu 85 90 95

Gly Ala Gln Val Gln Val Met Gly Thr Lys Gly Ala Thr Met Thr Ile 100 105 110

Leu Glu Phe Pro Glu Gly Thr Asp Gly Arg Leu Ile Val Arg Ser Glu 115 120 125

Asn Asp Thr Arg Arg Asn His Pro Ile Pro Pro Arg Gly Ser Leu Ser 130 135 140

Gln Cys Arg Ser Phe His His Gln Arg Cys Phe Asp Pro Val Ser His 145 150 155 160

Arg Pro Asp Arg Arg Leu Tyr Arg Cys Ala Gln Arg Arg Pro Pro Thr 165 170 175

Thr Asp His Arg Pro Arg Cys His Gln Ser Ser Glu Ser Ser Pro Trp 180 185 190

Cys Leu Arg Ile Ser Ser His Pro Pro Ala Gly Leu Phe Asp Leu Thr 195 200 205

Glu Ala Phe Lys Thr Ser Arg Gln Ile Gly Leu Ala Pro Leu Ser Ser 210 215 220

Leu Ser Thr Pro Pro Asp Gln Leu Val Arg Leu Ala Ala Ala Thr Gly 225 230 235 240

Phe Ser Phe Val Gly Leu Arg Val Ile Ala Val Thr Pro Asn Glu Arg 245 250 255

Val Tyr Asp Leu Ser Pro Gly Ser Pro Leu Leu Ala Ala Thr Gln Gln 260 265 270

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Val Asn Ala Asp Thr Thr Arg Glu Ala Trp Leu Pro Ala Leu Glu Ala 290 295 300

Ala Gly Ala Leu Gly Ala Lys Thr Phe Thr Ile Ala Ala Gly Asp Asp 305 310 315 320

Asn Ile Ala Pro Leu Thr Asp Thr Ile Gly Ala Met Val Asp Asp Ala 325 330 335

Arg Asp Phe Gly Val Thr Pro Ala Leu Glu Pro Ile Ser Tyr Arg Ser 340 345 350

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Ser	Ala	Val 35	Cys	Asp	Val	Asp	Gly 40	Ala	Lys	Ala	Ser	Glu 45	Thr	Ala	Ala	
Lys	Tyr 50.		Ile	Ser	Pro	Ser 55	Phe	Thr	Ser	Val	Asp 60	Glu	Ile	Leu	Ala	
Ser 65	Gly	Val	Asp	Ile	Val 70	Ala	Val	Cys	Thr	Pro 75	His	Pro	Thr	His	Glu 80	
Thr	Val	Val	Leu	Ala 85	Ala	Ala	Ala	Ala	Gly 90	Val	His	Val	Leu	Cys 95	Glu	

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					caa Gln											643
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Ser	Gly	Glu 195	Ile	Thr	Phe	Asp	Leu 200	Ser	Asp	Ala	Gln	Pro 205	Gly	Ser	Ala	
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caa cga ca Gln Arg Hi			Ile A								336
gtt gaa aac Val Glu Ly: 11:	Phe Pro										384
aca agt car Thr Ser Hi: 130			Lys \								432
tat gca ato Tyr Ala Ilo 145											480
ctt tat aa Leu Tyr Asi		Asp Ile			p His						528
aat aca gaq Asn Thr Glu			Gly 7								576
ttt aat tta Phe Asn Lei 195	Asp Asn										624
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ctt gga aag Leu Gly Lys 225											720
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Leu Gly Ala Val His Gly Leu Lys Tyr Trp Tyr Asn Tyr Thr Ser Asp 40 Asp Leu Ile Asn Phe Lys Pro Glu Gly Pro Ile Leu Asn Pro Asp Thr Lys Tyr Asp Ser His Gly Val Tyr Ser Gly Ser Ala Phe Glu Tyr Asn Gly His Leu Tyr Tyr Met Tyr Thr Gly Asn His Arg Asp Asn His Trp Gln Arg His Ala Ser Gln Met Ile Ala Arg Leu Lys Glu Asp Gly Ser Val Glu Lys Phe Pro Lys Pro Val Ile Ser Gln Gln Pro Glu Gly Tyr Thr Ser His Phe Arg Asp Pro Lys Val Phe Lys Tyr Gly Glu Lys Tyr 135 Tyr Ala Ile Ile Gly Ala Gln Asn Asn Asp Gln Gln Gly Arg Leu Leu 145 150 Leu Tyr Asn Thr Glu Asp Ile Ile Asn Trp His Tyr Leu Gly Glu Ile Asn Thr Glu Leu Asp Asp Phe Gly Tyr Met Trp Glu Cys Pro Asp Tyr 185 Phe Asn Leu Asp Asn Gln Asp Val Ile Leu Ile Cys Pro Gln Gly Ile 195 200 Glu Pro Lys Gly Asn Gln Phe Lys Asn Ile Tyr Gln Ser Gly Tyr Ile Leu Gly Lys Phe Asp Ile Glu Lys Leu Thr Tyr Glu His Glu Asn Phe 230 235

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<223> RXA02061

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Met Thr Asn Val Ser
1 5
ggg tat cac cga cca gag ctg cac atc acc gct gaa agt ggt gtt ttg 16

Gly	Tyr	His	Arg	Pro 10	Glu	Leu	His	Ile	Thr 15	Ala	Glu	Ser	Gly	Val 20	Leu	•
									gac Asp							211
									ggc Gly							259
									gat Asp							307
									cgc Arg							355
									acc Thr 95							403
									aac Asn							451
ctg Leu	atc Ile	aat Asn 120	gag Glu	gac Asp	gag Glu	ctg Leu	ggg Gly 125	ctc Leu	gat Asp	cca Pro	gat Asp	gtc Val 130	tcc Ser	cga Arg	atc Ile	499
									tat Tyr							547
tgc Cys 150	gtt Val	atc Ile	cca Pro	ggt Gly	tgg Trp 155	gaa Glu	gac Asp	caa Gln	gga Gly	aac Asn 160	cgc Arg	gat Asp	gaa Glu	ggc Gly	cac His 165	595
Cys 150 tca	Val gga	Ile tgg	Pro ttg	Gly	Trp 155 ctc	Glu gca	Asp gtt	Gln	gga Gly ggc Gly 175	Asn 160 cca	Arg gtt	Asp gaa	Glu gcc	Gly	His 165 aca	595 643
Cys 150 tca Ser	Val gga Gly gtg	Ile tgg Trp	Pro ttg Leu ctc	atg Met 170	Trp 155 ctc Leu tcg	Glu gca Ala cca	Asp gtt Val gat	Gln act Thr	Gly ggc Gly	Asn 160 cca Pro	Arg gtt Val	Asp gaa Glu tcc	Glu gcc Ala att	CCa Pro 180	His 165 aca Thr	
Cys 150 tca Ser gta Val	yal gga Gly gtg Val	tgg Trp gtc Val	ttg Leu ctc Leu 185	atg Met 170 gac Asp	Trp 155 ctc Leu tcg Ser	Glu gca Ala cca Pro	gtt Val gat Asp	Gln act Thr gga Gly 190 gga	ggc Gly 175	Asn 160 cca Pro gaa Glu	Arg gtt Val tgg Trp	gaa Glu tcc Ser	gcc Ala att Ile 195 gaa	CCa Pro 180 aca Thr	His 165 aca Thr ggt Gly	643
Cys 150 tca Ser gta Val ccc Pro	yal gga Gly gtg Val ctg Leu	tgg Trp gtc Val tct Ser 200	ttg Leu ctc Leu 185 ctc Leu	atg Met 170 gac Asp aac	Trp 155 ctc Leu tcg Ser ggc Gly	Glu gca Ala cca Pro ctc Leu	gtt Val gat Asp tct Ser 205	Gln act Thr gga Gly 190 gga Gly	ggc Gly 175 aga Arg	Asn 160 cca Pro gaa Glu gag Glu	gtt Val tgg Trp tca Ser	gaa Glu tcc Ser gac Asp 210	gcc Ala att Ile 195 gaa Glu	CCa Pro 180 aca Thr gtt Val	His 165 aca Thr ggt Gly cta Leu	643 691
Cys 150 tca Ser gta Val ccc Pro gtt Val	yal gga Gly gtg Val ctg Leu gct Ala 215	tct Ser 200 cct Pro	ttg Leu ctc Leu 185 ctc Leu cgc Arg	atg Met 170 gac Asp aac Asn atg Met	Trp 155 ctc Leu tcg Ser ggc Gly att Ile	Glu gca Ala cca Pro ctc Leu cgt Arg 220 acc	gtt Val gat Asp tct Ser 205 ctg Leu	Gln act Thr gga Gly 190 gga Gly cgc Arg	ggc Gly 175 aga Arg tta Leu	Asn 160 cca Pro gaa Glu gag Glu gaa Glu	gtt Val tgg Trp tca Ser gtg Val 225	gaa Glu tcc Ser gac Asp 210 gat Asp	gcc Ala att Ile 195 gaa Glu cat His	CCa Pro 180 aca Thr gtt Val gaa Glu	His 165 aca Thr ggt Gly cta Leu atc Ile	643 691 739

250 255 260 cca ttt acc cgc atc gat ttt ggc cat gat ttc tct cgc ccc cgc aac 931 Pro Phe Thr Arg Ile Asp Phe Gly His Asp Phe Ser Arg Pro Arg Asn 265 270 acc aac tac qcc qaa acc acc atc qqc tac qac ttc qcc cac atc ttt 979 Thr Asn Tyr Ala Glu Thr Thr Ile Gly Tyr Asp Phe Ala His Ile Phe 280 ggt ctc atg aat ggc gta ggt cgt ttg gac tcc ccc act gag cat ctc 1027 Gly Leu Met Asn Gly Val Gly Arg Leu Asp Ser Pro Thr Glu His Leu 295 300 agt tgg aag gaa gaa ggc tgg gca aac gct att tct ttc cca cgt att 1075 Ser Trp Lys Glu Glu Gly Trp Ala Asn Ala Ile Ser Phe Pro Arg Ile 310 315 gtc acg ctc cag gac ggt acg gtc ttc cag acc cct cca gaa gga ttg 1123 Val Thr Leu Gln Asp Gly Thr Val Phe Gln Thr Pro Pro Glu Gly Leu 330 335 ctt gat gcc att cat gaa tcc gag gca gcg gca ggt tgg acc gga ctg 1171 Leu Asp Ala Ile His Glu Ser Glu Ala Ala Ala Gly Trp Thr Gly Leu tgc gaa atc cca tca aac agc gca gtt gaa gtg gcg ttg aag gac caa 1219 Cys Glu Ile Pro Ser Asn Ser Ala Val Glu Val Ala Leu Lys Asp Gln 365 gaa ggt gaa atc gct gca aca atc act cac cgc cac aat cag cta gtc 1267 Glu Gly Glu Ile Ala Ala Thr Ile Thr His Arg His Asn Gln Leu Val 380 gtt gat cgg tcc atg aac ccc aac cac gcg ggt gat cca cac gcg att 1315 Val Asp Arg Ser Met Asn Pro Asn His Ala Gly Asp Pro His Ala Ile 395 400 gca cca ttg act gat gat gaa aca gat tca ctg ttc att gtc gtt gac 1363 Ala Pro Leu Thr Asp Asp Glu Thr Asp Ser Leu Phe Ile Val Val Asp 410 415 ggc tct aca gta gaa gtt ttt gct gat ggc ggt tat gta tca atg gca 1411 Gly Ser Thr Val Glu Val Phe Ala Asp Gly Gly Tyr Val Ser Met Ala 425 age egt gtg tat tte aac aac gga eea tte age gaa ttt gag gte ace 1459 Ser Arg Val Tyr Phe Asn Asn Gly Pro Phe Ser Glu Phe Glu Val Thr 445 acc acc ggt gac gca agc att att cgc cag gaa agt cac ttc cct gtt 1507 Thr Thr Gly Asp Ala Ser Ile Ile Arg Gln Glu Ser His Phe Pro Val 455 460 gat ttc agt tcg gtg tcc cta gat ata gat gat ctc act gcg ctc atg 1555 Asp Phe Ser Ser Val Ser Leu Asp Ile Asp Asp Leu Thr Ala Leu Met 470 475 480 cag ttc gat gaa aac gaa ccg cat gaa ggc cca gtg aga taagagttag 1604 Gln Phe Asp Glu Asn Glu Pro His Glu Gly Pro Val Arg

490

atgcgttcca gcc 1617

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<211> 498

<212> PRT

<213> Corynebacterium glutamicum

<400> 352

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Thr Trp His Phe Phe His Gln Tyr Arg Pro Ser Pro Asp His Gly Pro 35 40 45

Arg Trp Ala His Gln Phe Ala Glu Arg Thr Pro Phe Val Trp Asp Ile 50 55 60

Cys Asp Asp Val Leu Ala Pro Glu Gly Asp Glu Thr Gln Val Arg Ala 65 70 75 80

Gly Ser Val Val Ser Asn Asn Gly Gly Val Asp Leu Tyr Phe Thr Ser 85 90 95

Val Val Gly Pro Thr Ser Thr Ile Gln Leu Ala His Ile Asn Asn Ile 100 105 110

Arg Gly Thr Thr Glu Leu Ile Asn Glu Asp Glu Leu Gly Leu Asp Pro 115 120 125 .

Asp Val Ser Arg Ile Gly Glu Val Val Gly Asn Thr Asp Gly Tyr Val 130 135 140

Lys Phe Arg Ser Pro Cys Val Ile Pro Gly Trp Glu Asp Gln Gly Asn 145 150 155 160

Arg Asp Glu Gly His Ser Gly Trp Leu Met Leu Ala Val Thr Gly Pro 165 170 175

Val Glu Ala Pro Thr Val Val Leu Asp Ser Pro Asp Gly Arg Glu 180 185 190

Trp Ser Ile Thr Gly Pro Leu Ser Leu Asn Gly Leu Ser Gly Leu Glu
195 200 205

Ser Asp Glu Val Leu Val Ala Pro Arg Met Ile Arg Leu Arg Asp Glu 210 215 220

Val Asp His Glu Ile Tyr Asp Val Leu Ile Val Thr Ile Glu Gln Asp 225 230 235 240

Gly Ile Asp Ile Ser Gly Tyr Leu Val Gly Gln Leu Asn Gly Ser Glu 245 250 255

Phe Asp Val Lys Thr Pro Phe Thr Arg Ile Asp Phe Gly His Asp Phe 260 265 270

Ser Arg Pro Arg Asn Thr Asn Tyr Ala Glu Thr Thr Ile Gly Tyr Asp Phe Ala His Ile Phe Gly Leu Met Asn Gly Val Gly Arg Leu Asp Ser 295 Pro Thr Glu His Leu Ser Trp Lys Glu Glu Gly Trp Ala Asn Ala Ile Ser Phe Pro Arg Ile Val Thr Leu Gln Asp Gly Thr Val Phe Gln Thr Pro Pro Glu Gly Leu Leu Asp Ala Ile His Glu Ser Glu Ala Ala Ala Gly Trp Thr Gly Leu Cys Glu Ile Pro Ser Asn Ser Ala Val Glu Val 360 Ala Leu Lys Asp Gln Glu Gly Glu Ile Ala Ala Thr Ile Thr His Arg 375 His Asn Gln Leu Val Val Asp Arg Ser Met Asn Pro Asn His Ala Gly 385 390 395 Asp Pro His Ala Ile Ala Pro Leu Thr Asp Asp Glu Thr Asp Ser Leu Phe Ile Val Val Asp Gly Ser Thr Val Glu Val Phe Ala Asp Gly Gly 425 Tyr Val Ser Met Ala Ser Arg Val Tyr Phe Asn Asn Gly Pro Phe Ser Glu Phe Glu Val Thr Thr Gly Asp Ala Ser Ile Ile Arg Gln Glu 455 Ser His Phe Pro Val Asp Phe Ser Ser Val Ser Leu Asp Ile Asp Asp 470 475 Leu Thr Ala Leu Met Gln Phe Asp Glu Asn Glu Pro His Glu Gly Pro 485 490 Val Arg

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ggc tca ctq cqc acc tac cca tgg ggt tca aga aca ctg atc gct gat 163 Gly Ser Leu Arg Thr Tyr Pro Trp Gly Ser Arg Thr Leu Ile Ala Asp ctc aaa ggc gaa gaa tca cca tcg tct cgc cca gag gcc gaa gtc tgg 211 Leu Lys Gly Glu Glu Ser Pro Ser Ser Arg Pro Glu Ala Glu Val Trp 30 ttc qqt qcc cac cca qqa tca cca tca acc atc ggt gga aac gca ctc Phe Gly Ala His Pro Gly Ser Pro Ser Thr Ile Gly Gly Asn Ala Leu 307 aac gaa gtc atc gca gcg aac ccc gaa gaa gca ttg ggc acg cgt gtt Asn Glu Val Ile Ala Ala Asn Pro Glu Glu Ala Leu Gly Thr Arg Val 55 60 qcc qaa qcq ttt qaa aat qaq ctt cca ttc ctc ctc aaa atc ctc gca Ala Glu Ala Phe Glu Asn Glu Leu Pro Phe Leu Leu Lys Ile Leu Ala gcg gga gca ccc cta tca ctg cag gcc cac cca tcg ctg gaa cag gcc Ala Gly Ala Pro Leu Ser Leu Gln Ala His Pro Ser Leu Glu Gln Ala cqt qaa qqa ttc qcc cqc qaa aac tca gca gga att gac ctc ggc gca Arg Glu Gly Phe Ala Arg Glu Asn Ser Ala Gly Ile Asp Leu Gly Ala 105 ccg aac cgc aac tac cqc gac cca aac cac aag cca gag ctg atc gtt 499 Pro Asn Arg Asn Tyr Arg Asp Pro Asn His Lys Pro Glu Leu Ile Val 120 125 547 get etc aeg gaa tte atc geg atg gea gge tte ege eea etg egg aac Ala Leu Thr Glu Phe Ile Ala Met Ala Gly Phe Arg Pro Leu Arg Asn 135 140 acc ctc acc att ttc gac gcc ctc gcc tgc gaa ccc ctc gac cgc tac Thr Leu Thr Ile Phe Asp Ala Leu Ala Cys Glu Pro Leu Asp Arg Tyr 155 cgc agc atg ctc acc gtc gac aac gag gaa gaa tcc ctc cgc gca ctg Arg Ser Met Leu Thr Val Asp Asn Glu Glu Glu Ser Leu Arg Ala Leu 170 175 ttt acc acc tgg atc acc atc ccc atc ggt aaa cga cac gaa ctc atc Phe Thr Thr Trp Ile Thr Ile Pro Ile Gly Lys Arg His Glu Leu Ile 190 gat gcc ctc atc agc aac gcc cac acc tac ctt gag gca agc gat cgt 739 Asp Ala Leu Ile Ser Asn Ala His Thr Tyr Leu Glu Ala Ser Asp Arg 787 gac gag gac atc gca ttc gtg ctc tca cac atc atc gag ctc aac gaa Asp Glu Asp Ile Ala Phe Val Leu Ser His Ile Ile Glu Leu Asn Glu 215 220 cag tac ecc qqc gat qtc qqc gtt etg ggt get etg etg ttg aac tte 835 Gln Tyr Pro Gly Asp Val Gly Val Leu Gly Ala Leu Leu Leu Asn Phe 230 240

	aaa Lys															883
	gca Ala															931
	gtg Val															979
	gtg Val 295															1027
	gaa Glu															1075
	caa Gln															1123
gat Asp	ggt Gly	ccc Pro	atg Met 345	att Ile	gtt Val	ctg Leu	tgc Cys	acc Thr 350	tcc Ser	gga Gly	act Thr	gtt Val	tcc Ser 355	ttg Leu	gaa Glu	1171
	ggg Gly															1219
	gca Ala 375															1267
	ctc Leu			_	taga	tctt	tt t	agat	taaa	a to	a					1305
)> 35 L> 39															

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<213> Corynebacterium glutamicum

<400> 354

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Thr Leu Ile Ala Asp Leu Lys Gly Glu Glu Ser Pro Ser Ser Arg Pro 25

Glu Ala Glu Val Trp Phe Gly Ala His Pro Gly Ser Pro Ser Thr Ile

Gly Gly Asn Ala Leu Asn Glu Val Ile Ala Ala Asn Pro Glu Glu Ala

Leu Gly Thr Arg Val Ala Glu Ala Phe Glu Asn Glu Leu Pro Phe Leu

65					70					75					80
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Ser	Leu	Glu	Gln 100	Ala	Arg	Glu	Gly	Phe 105	Ala	Arg	Glu	Asn	Ser 110	Ala	Gly
Ile	Asp	Leu 115	Gly	Ala	Pro	Asn	Arg 120	Asn	Tyr	Arg	Asp	Pro 125	Asn	His	Lys
Pro	Glu 130	Leu	Ile	Val	Ala	Leu 135	Thr	Glu	Phe	Ile	Ala 140	Met	Ala	Gly	Phe
Arg 145	Pro	Leu	Arg	Asn	Thr 150	Leu	Thr	Ile	Phe	Asp 155	Ala	Leu	Ala	Cys	Glu 160
Pro	Leu	Asp	Arg	Tyr 165	Arg	Ser	Met	Leu	Thr 170	Val	Asp	Asn	Glu	Glu 175	Glu
Ser	Leu	Arg	Ala 180	Leu	Phe	Thr	Thr	Trp 185	Ile	Thr	Ile	Pro	Ile 190	Gly	Lys
Arg	His	Glu 195	Leu	Ile	Asp	Ala	Leu 200	Ile	Ser	Asn	Ala	His 205	Thr	Tyr	Leu
Glu	Ala 210	Ser	Asp	Arg	Asp	Glu 215	Asp	Ile	Ala	Phe	Val 220	Leu	Ser	His	Ile
Ile 225	Glu	Leu	Asn	Glu	Gln 230	Tyr	Pro	Gly	Asp	Val 235	Gly	Val	Leu	Gly	Ala 240
Leu	Leu	Leu	Asn	Phe 245	Tyr	Lys	Leu	Ala	Pro 250	Gly	Glu	Ala	Leu	Tyr 255	Leu
Asp	Ala	Ala	Asn 260	Leu	His	Ala	Tyr	Ile 265	Ser	Gly	Leu	Gly	Val 270	Glu	Ile
Met	Ala	Asn 275	Ser	Asp	Asn	Val	Leu 280	Arg	Gly	Gly	Leu	Thr 285	Ser	Lys	Tyr
Val	Asp 290	Val	Pro	Glu	Leu	Val 295	Arg	Val	Leu	Asp	Phe 300	Asn	Ser	Leu	Glu
Asn 305	Ala	Arg	Val	Asp	Val 310	Glu	Glu	Asp	Gly	Ala 315	Thr	Thr	His	Tyr	Pro 320
Val	Pro	Ile	Asn	Glu 325	Phe	Gln	Leu	Asp	Arg 330	Val	Ala	Val	Gln	Gly 335	Glu
Ala	Glu	Ala	Asn 340	His	Asp	Gly	Pro	Met 345	Ile	Val	Leu	Cys	Thr 350	Ser	Gly
Thr	Val	Ser 355	Leu	Glu	Ala	Gly	Glu 360	Lys	Thr	Leu	Glu	Val 365	Ala	Ala	Gly
His	Ala 370	Ala	Trp	Val	Pro	Ala 375	Asn	Asp	Pro	Thr	Ile 380	Ala	Met	Arg	Ser
Glu 385	Asp	Ala	Glu	Val	Phe 390	Leu	Ala	Arg	Val						

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aac Asn	ttc Phe	tac Tyr	aaa Lys 20	ctt Leu	gcc Ala	cca Pro	ggc Gly	gaa Glu 25	gcc Ala	ctc Leu	tac Tyr	ctc Leu	gac Asp 30	gcc Ala	gca Ala	96
	ctt Leu															144
	gac Asp 50															192
ccg Pro 65	gag Glu	ctt Leu	gtg Val	cgc Arg	gtg Val 70	ttg Leu	gat Asp	ttc Phe	aac Asn	tct Ser 75	ttg Leu	gaa Glu	aac Asn	gct Ala	cgc Arg 80	240
	gac Asp															288
aac Asn	gaa Glu	ttc Phe	caa Gln 100	ctc Leu	gat Asp	cgc Arg	gtt Val	gca Ala 105	gtt Val	cag Gln	ggc Gly	gaa Glu	gca Ala 110	gaa Glu	gcc Ala	336
aac Asn	cac His	gat Asp 115	ggt Gly	ccc Pro	atg Met	att Ile	gtt Val 120	ctg Leu	tgc Cys	acc Thr	tcc Ser	gga Gly 125	act Thr	gtt Val	tcc Ser	384
ttg Leu	gaa Glu 130	gca Ala	Gly ggg	gag Glu	aag Lys	acc Thr 135	ctc Leu	gaa Glu	gta Val	gca Ala	gca Ala 140	ggt Gly	cac His	gcc Ala	gca Ala	432
tgg Trp 145	gtt Val	cca Pro	gca Ala	aac Asn	gac Asp 150	cca Pro	acc Thr	att Ile	gcg Ala	atg Met 155	cgt Arg	tct Ser	gag Glu	gac Asp	gca Ala 160	480
gaa Glu	gta Val	ttc Phe	ctc Leu	gct Ala 165	agg Arg	gtt Val	taga	itctt	tt t	agat	taaa	a to	a			524

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<400> 356

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Asn Phe Tyr Lys Leu Ala Pro Gly Glu Ala Leu Tyr Leu Asp Ala Ala 20 25 30

Asn Leu His Ala Tyr Ile Ser Gly Leu Gly Val Glu Ile Met Ala Asn 35 40 45

Ser Asp Asn Val Leu Arg Gly Gly Leu Thr Ser Lys Tyr Val Asp Val 50 55 60

Pro Glu Leu Val Arg Val Leu Asp Phe Asn Ser Leu Glu Asn Ala Arg 65 70 75 80

Val Asp Val Glu Glu Asp Gly Ala Thr Thr His Tyr Pro Val Pro Ile 85 90 95

Asn Glu Phe Gln Leu Asp Arg Val Ala Val Gln Gly Glu Ala Glu Ala 100 105 110

Asn His Asp Gly Pro Met Ile Val Leu Cys Thr Ser Gly Thr Val Ser 115 120 125

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Trp Val Pro Ala Asn Asp Pro Thr Ile Ala Met Arg Ser Glu Asp Ala 145 150 155 160

Glu Val Phe Leu Ala Arg Val

<210> 357

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<212> DNA

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<220>

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<222> (101)..(808)

<223> FRXA01373

<400> 357

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Met Glu Leu Glu

1 5

ggc tca ctg cgc acc tac cca tgg ggt tca aga aca ctg atc gct gat 163 Gly Ser Leu Arg Thr Tyr Pro Trp Gly Ser Arg Thr Leu Ile Ala Asp

ctc aaa ggc gaa gaa tca cca tcg tct cgc cca gag gcc gaa gtc tgg 211 Leu Lys Gly Glu Glu Ser Pro Ser Ser Arg Pro Glu Ala Glu Val Trp 25 30 35

		-					cca Pro 45							-		259
		-		-			ccc Pro	-	-	_	-		-	-	-	307
							ctt Leu									355
							cag Gln									403
				_	_	_	aac Asn		-			-			-	451
							cca Pro 125									499
							atg Met									547
							ctc Leu									595
							aac Asn									643
							ccc Pro				_		-			691
							cac His 205									739
							ctc Leu									787
				gat Asp												808
<211 <212)> 35 .> 23 !> PR !> Co	86 RT	ebact	eriu	um gl	utam	nicum	ı								
)> 35		Ten	C1	Cl	Co~	Ton	A = ~	ሞ⊳∽	Tur	Dro	T	C1	Co-	λ = σ	

Met Glu Leu Glu Gly Ser Leu Arg Thr Tyr Pro Trp Gly Ser Arg

10

15

5

Thr Leu Ile Ala Asp Leu Lys Gly Glu Glu Ser Pro Ser Ser Arg Pro 25 Glu Ala Glu Val Trp Phe Gly Ala His Pro Gly Ser Pro Ser Thr Ile 40 Gly Gly Asn Ala Leu Asn Glu Val Ile Ala Ala Asn Pro Glu Glu Ala Leu Gly Thr Arg Val Ala Glu Ala Phe Glu Asn Glu Leu Pro Phe Leu Leu Lys Ile Leu Ala Ala Gly Ala Pro Leu Ser Leu Gln Ala His Pro 90 Ser Leu Glu Gln Ala Arg Glu Gly Phe Ala Arg Glu Asn Ser Ala Gly Ile Asp Leu Gly Ala Pro Asn Arg Asn Tyr Arg Asp Pro Asn His Lys Pro Glu Leu Ile Val Ala Leu Thr Glu Phe Ile Ala Met Ala Gly Phe 130 135 Arg Pro Leu Arg Asn Thr Leu Thr Ile Phe Asp Ala Leu Ala Cys Glu 150 Pro Leu Asp Arg Tyr Arg Ser Met Leu Thr Val Asp Asn Glu Glu Glu Ser Leu Arg Ala Leu Phe Thr Thr Trp Ile Thr Ile Pro Ile Gly Lys 185 Arg His Glu Leu Ile Asp Ala Leu Ile Ser Asn Ala His Thr Tyr Leu 200 Glu Ala Ser Asp Arg Asp Glu Asp Ile Ala Phe Val Leu Ser His Ile 210 215 Ile Glu Leu Asn Glu Gln Tyr Pro Gly Asp Val Gly 230 <210> 359 <211> 1775 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (1)..(1752) <223> RXA02611 <400> 359 gat gcg tgg tcg gat cct atg gct acg tgg cgt cat gcg att acc act 48 Asp Ala Trp Ser Asp Pro Met Ala Thr Trp Arg His Ala Ile Thr Thr 10 aag att gag gcc ggc cag ggt tcg gat gag ttg tat aac gac ttt gag

Ly	3 Ile	Glu	Ala 20	Gly	Gln	Gly	Ser	Asp 25	Glu	Leu	Tyr	Asn	Asp 30	Phe	Glu	
	ggg Gly															144
	agg Arg 50															192
	gta Val															240
	tta Leu															288
	g cag 1 Gln															336
	g ctt Leu															384
gti Va	cat His 130	ggc Gly	act Thr	ttc Phe	gct Ala	acc Thr 135	act Thr	gct Ala	cag Gln	gcg Ala	ttg Leu 140	gag Glu	cgt Arg	gtc Val	gcg Ala	432
	atg Met															480
	gtc Val															528
gat Asp	gtg Val	ggt Gly	tcg Ser 180	ccg Pro	tgg Trp	gct Ala	att Ile	ggt Gly 185	tct Ser	aaa Lys	gat Asp	ggt Gly	ggg Gly 190	cat His	gat Asp	576
gca Ala	acg Thr	cat His 195	ccg Pro	cgg Arg	ttg Leu	ggc Gly	acc Thr 200	att Ile	gaa Glu	gat Asp	ttc Phe	cag Gln 205	gcg Ala	ttg Leu	ttg Leu	624
gct Ala	cgc Arg 210	gca Ala	cgg Arg	gaa Glu	ctc Leu	aat Asn 215	ttg Leu	gaa Glu	gtt Val	gca Ala	ctc Leu 220	gat Asp	cta Leu	gct Ala	ctg Leu	672
	gct Ala															720
	gtg Val															768
	tac Tyr															816

270 260 265 atc tac gaa gag gtc tat cgt gtg gtg aag ttc tgg gtg gat ttg ggt 864 Ile Tyr Glu Glu Val Tyr Arg Val Val Lys Phe Trp Val Asp Leu Gly 280 gtg acc aca ttc cgc gtg gat aac ccg cac act aag ccc gct aat ttc 912 Val Thr Thr Phe Arg Val Asp Asn Pro His Thr Lys Pro Ala Asn Phe 295 290 tgg cag tgg ctt att tct gcc atc cat aaa tca aac cct gag gtc att 960 Trp Gln Trp Leu Ile Ser Ala Ile His Lys Ser Asn Pro Glu Val Ile 315 ttc cta gcg gag gcg tct act cgc ccg gca cgt ctg tat ttc ttg tcc 1008 Phe Leu Ala Glu Ala Ser Thr Arg Pro Ala Arg Leu Tyr Phe Leu Ser 330 325 1056 aag att ggt ttc tcc cag tct tac acc tac ttc acc tgg aag gtc acc Lys Ile Gly Phe Ser Gln Ser Tyr Thr Tyr Phe Thr Trp Lys Val Thr 340 345 aac gag gag etc acc gag ttc gct act gag atc gcc ccc atg gcg gat 1104 Asn Glu Glu Leu Thr Glu Phe Ala Thr Glu Ile Ala Pro Met Ala Asp 355 360 1152 att tot ogt oog aac otg ttt gtg aac act ooc gac att ttg cat gog Ile Ser Arg Pro Asn Leu Phe Val Asn Thr Pro Asp Ile Leu His Ala 380 370 375 tct ctg cag cat ggt gga cgc gcc atg ttc gct atc cgc gcc gca ttg 1200 Ser Leu Gln His Gly Gly Arg Ala Met Phe Ala Ile Arg Ala Ala Leu 385 390 1248 gcc gcc acg atg tct cct gtg tgg ggc gta tat tcc gga tat gag ctc Ala Ala Thr Met Ser Pro Val Trp Gly Val Tyr Ser Gly Tyr Glu Leu 405 ttt gag cac gag gcc gtc aag cct ggt tcg gaa gag tac ttg gat tct 1296 Phe Glu His Glu Ala Val Lys Pro Gly Ser Glu Glu Tyr Leu Asp Ser 425 gag aag tac gag ctg cgt ccc cgc gat ttc gag ggt gct ctg gaa cgt 1344 Glu Lys Tyr Glu Leu Arg Pro Arg Asp Phe Glu Gly Ala Leu Glu Arg 435 ggc gat tot otc gag gat tac atc gct ctg ctc aac cag atc cgt cgc 1392 Gly Asp Ser Leu Glu Asp Tyr Ile Ala Leu Leu Asn Gln Ile Arg Arg gcg aac cct gcc ttg cag caa cta cgc aac atc cac ttc cac gaa gcg 1440 Ala Asn Pro Ala Leu Gln Gln Leu Arg Asn Ile His Phe His Glu Ala 475 1488 gac aat gat cag atc atc gcc tac tcc aag gtt gat gct ttg acc gga Asp Asn Asp Gln Ile Ile Ala Tyr Ser Lys Val Asp Ala Leu Thr Gly aat acc gtg ttg att gtg gtc aac ttg gat cca cgt agt gct cgt gag 1536 Asn Thr Val Leu Ile Val Val Asn Leu Asp Pro Arg Ser Ala Arg Glu

505

500

							gga Gly 520	Ala		Gly						1584
							atc Ile								tca Ser	1632
							gag Glu									1680
ttt Phe	gtt Val	ctt Leu	cct Pro	gaa Glu 565	ctt Leu	cca Pro	gcg Ala	tct Ser	cgc Arg 570	cgt Arg	gag Glu	cgt Arg	ctc Leu	gcg Ala 575	tgg Trp	1728
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<212> PRT

<213> Corynebacterium glutamicum

<400> 360

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Lys Ile Glu Ala Gly Gln Gly Ser Asp Glu Leu Tyr Asn Asp Phe Glu 20 25 30

His Gly Ala Gln Leu Phe Glu Arg Ala Ala Glu Asn Leu Ser Lys Glu 35 40 45

Asp Arg Thr Ala Leu Phe Asp Val Ala Ser Ser Leu Arg Arg Gly Gly 50 55 60

Asp Val Arg Ala Arg Leu Ala Pro Ala Leu Thr Ala Ser Val Thr His 65 70 75 80

Leu Leu Glu Leu Asn Pro Leu Arg Glu Leu Val Thr Met Gly Glu Asn 85 90 95

Leu Gln Val Arg Val Glu Arg Arg Ala Ala Leu Val Asn Ser Trp Tyr
100 105 110

Glu Leu Phe Pro Arg Ser Thr Gly Gly Trp Asp Glu Ser Gly Thr Pro 115 120 125

Val His Gly Thr Phe Ala Thr Thr Ala Gln Ala Leu Glu Arg Val Ala 130 135 140

Lys Met Gly Phe Asp Thr Val Tyr Phe Pro Pro Ile His Pro Ile Gly 145 150 155 160

Glu Val Asn Arg Lys Gly Arg Asn Asn Thr Leu Thr Pro Glu Pro His 165 170 175

Asp Val Gly Ser Pro Trp Ala Ile Gly Ser Lys Asp Gly Gly His Asp Ala Thr His Pro Arg Leu Gly Thr Ile Glu Asp Phe Gln Ala Leu Leu Ala Arg Ala Arg Glu Leu Asn Leu Glu Val Ala Leu Asp Leu Ala Leu 215 Gln Ala Ala Pro Asp His Pro Trp Ala Gln Glu His Arg Glu Phe Phe 235 Thr Val Leu Ala Asp Gly Thr Ile Ala Tyr Ala Glu Asn Pro Pro Lys 250 Lys Tyr Gln Asp Ile Tyr Pro Ile Asn Phe Asp Asn Asp Ala Pro Lys 265 Ile Tyr Glu Glu Val Tyr Arg Val Val Lys Phe Trp Val Asp Leu Gly Val Thr Thr Phe Arg Val Asp Asn Pro His Thr Lys Pro Ala Asn Phe 295 Trp Gln Trp Leu Ile Ser Ala Ile His Lys Ser Asn Pro Glu Val Ile 315 310 Phe Leu Ala Glu Ala Ser Thr Arg Pro Ala Arg Leu Tyr Phe Leu Ser 330 Lys Ile Gly Phe Ser Gln Ser Tyr Thr Tyr Phe Thr Trp Lys Val Thr 345 Asn Glu Glu Leu Thr Glu Phe Ala Thr Glu Ile Ala Pro Met Ala Asp 360 Ile Ser Arg Pro Asn Leu Phe Val Asn Thr Pro Asp Ile Leu His Ala Ser Leu Gln His Gly Gly Arg Ala Met Phe Ala Ile Arg Ala Ala Leu 390 395 Ala Ala Thr Met Ser Pro Val Trp Gly Val Tyr Ser Gly Tyr Glu Leu 405 410 Phe Glu His Glu Ala Val Lys Pro Gly Ser Glu Glu Tyr Leu Asp Ser 425 Glu Lys Tyr Glu Leu Arg Pro Arg Asp Phe Glu Gly Ala Leu Glu Arg Gly Asp Ser Leu Glu Asp Tyr Ile Ala Leu Leu Asn Gln Ile Arg Arg Ala Asn Pro Ala Leu Gln Gln Leu Arg Asn Ile His Phe His Glu Ala 470 475 465 Asp Asn Asp Gln Ile Ile Ala Tyr Ser Lys Val Asp Ala Leu Thr Gly 490 Asn Thr Val Leu Ile Val Val Asn Leu Asp Pro Arg Ser Ala Arg Glu

			500					505					510			
Ala	Thr	Val 515	Arg	Leu	Asp	Leu	Gly 520	Ala	Leu	Gly	Leu	Glu 525	Ala	Gly	Ala	
Gln	Phe 530		Val	Arg	Asp	Ala 535	Ile	Thr	Gly	Ser	Arg 540	Tyr	Leu	Trp	Ser	
Glu 545	Thr	Asn	Phe	Val	Arg 550	Leu	Glu	Pro	Leu	Arg 555	Asp	Val	Ala	His	Ile 560	
Phe	.Val	Leu	Pro	Glu 565	Leu	Pro	Ala	Ser	Arg 570	Arg	Glu	Arg	Leu	Ala 575	Trp	
Arg	Glu	Ile	Lys 580	Thr	Tyr	Arg	Ala									
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taa	tttc	cca t	ctct	gtad	cc tt	ctat	caaq	g gat	ttato	catc	_		gtt Val	-		115
gcg	agc Ser	cac	atc	acc	atc	cct	gaa	gca	gat	ctg	Met 1 gcc	Thr	Val ctg	Asp	Pro 5 cac	115
gcg Ala tgc	agc	cac His	atc Ile cac	acc Thr 10 gat	atc Ile cct	cct Pro	gaa Glu gga	gca Ala ttt	gat Asp 15	ctg Leu ggt	Met 1 gcc Ala tgg	Thr cgc Arg	Val ctg Leu	Asp cgc Arg 20 acc	Pro 5 cac His	
gcg Ala tgc Cys	agc Ser	cac His cat His	atc Ile cac His 25	acc Thr 10 gat Asp	atc Ile cct Pro	cct Pro cat His	gaa Glu gga Gly	gca Ala ttt Phe 30	gat Asp 15 tat Tyr	ctg Leu ggt Gly	Met 1 gcc Ala tgg Trp	Thr cgc Arg cat His	Val ctg Leu gag Glu 35	Asp cgc Arg 20 acc Thr	Pro 5 cac His gaa Glu	163
gcg Ala tgc Cys gct Ala	agc Ser aac Asn	cac His cat His tcg Ser 40	atc Ile cac His 25 gtt Val	acc Thr 10 gat Asp atc Ile	atc Ile cct Pro cgc Arg	cct Pro cat His acg Thr	gaa Glu gga Gly cgc Arg 45	gca Ala ttt Phe 30 cag Gln	gat Asp 15 tat Tyr gtc Val	ctg Leu ggt Gly ggc Gly	Met 1 gcc Ala tgg Trp gcg Ala	Thr cgc Arg cat His acg Thr 50 atc	Val ctg Leu gag Glu 35 cag Gln	Asp cgc Arg 20 acc Thr gtt Val	Pro 5 cac His gaa Glu aat Asn	163 211
gcg Ala tgc Cys gct Ala ttg Leu	agc Ser aac Asn ggt Gly	cac His cat His tcg Ser 40 atc Ile	atc Ile cac His 25 gtt Val gac Asp	acc Thr 10 gat Asp atc Ile gac Asp	atc Ile cct Pro cgc Arg acc Thr	cct Pro cat His acg Thr tcc Ser 60	gaa Glu gga Gly cgc Arg 45 cac His	gca Ala ttt Phe 30 cag Gln gtc Val	gat Asp 15 tat Tyr gtc Val atg Met	ctg Leu ggt Gly ggc Gly acc Thr	Met 1 gcc Ala tgg Trp gcg Ala cct Pro 65	Thr cgc Arg cat His acg Thr 50 atc Ile	Val ctg Leu gag Glu 35 cag Gln ggc Gly	Asp cgc Arg 20 acc Thr gtt Val gac Asp	Pro 5 cac His gaa Glu aat Asn gac Asp	163 211 259

451

tac ttc ctc ccc acc gta ggc gag atg gat att tac ctc ttc tct gag Tyr Phe Leu Pro Thr Val Gly Glu Met Asp Ile Tyr Leu Phe Ser Glu

110 115 105 gga cgc cat gag cgt ttg tgg gag att ctc ggt gcc aac atc aag acc 499 Gly Arg His Glu Arg Leu Trp Glu Ile Leu Gly Ala Asn Ile Lys Thr 120 125 547 tac caa act gcg ctc gga aca gtt cgt ggc acc gca ttt act gtg tgg Tyr Gln Thr Ala Leu Gly Thr Val Arg Gly Thr Ala Phe Thr Val Trp 140 gct cca aac gca att ggc tgc gca gtg gtc ggt ggc ttc aac ggt tgg Ala Pro Asn Ala Ile Gly Cys Ala Val Val Gly Gly Phe Asn Gly Trp 150 155 aat gca tcc cag cat ccg atg cgt tct atg ggt ggt tcg ggt ctg tgg 643 Asn Ala Ser Gln His Pro Met Arg Ser Met Gly Gly Ser Gly Leu Trp 170 175 gag ctg ttc atc cca ggc ata gag gaa ggc gaa gtg tac aaa ttc gcc Glu Leu Phe Ile Pro Gly Ile Glu Glu Glu Glu Val Tyr Lys Phe Ala 190 185 739 gtc caa acc agg gaa ggc caa cgt cgt gat aag gcc gat ccg atg gct Val Gln Thr Arg Glu Gly Gln Arg Arg Asp Lys Ala Asp Pro Met Ala 200 cgt cgc gca gaa ctg gcg ccg gca acc gga tct att gtc gct tcc tct 787 Arg Arg Ala Glu Leu Ala Pro Ala Thr Gly Ser Ile Val Ala Ser Ser 215 220 835 gag tac cag tgg cag gat tcc gag tgg ctg cgc gag cgt tcc caa act Glu Tyr Gln Trp Gln Asp Ser Glu Trp Leu Arg Glu Arg Ser Gln Thr 230 . 235 883 gat etc gea tec aag eea atg agt gte tac gag gte eac etc ggt tet Asp Leu Ala Ser Lys Pro Met Ser Val Tyr Glu Val His Leu Gly Ser 250 tgg cgc tgg ggt aag aac tat gag gat ttg gct act gag ctg gtt gat Trp Arg Trp Gly Lys Asn Tyr Glu Asp Leu Ala Thr Glu Leu Val Asp 979 tac gtc gca gat ctt ggc tac acc cac gtg gaa ttc ctc cct gtc gca Tyr Val Ala Asp Leu Gly Tyr Thr His Val Glu Phe Leu Pro Val Ala 285 gag cac ccc ttc ggt ggt tcc tgg ggt tac cag gtc acc ggc tac tac 1027 Glu His Pro Phe Gly Gly Ser Trp Gly Tyr Gln Val Thr Gly Tyr Tyr 300 1075 gca ccg acc tct cgt tgg ggt act cca gat cag ttc cgt gcg cta gtc Ala Pro Thr Ser Arg Trp Gly Thr Pro Asp Gln Phe Arg Ala Leu Val 310 gac gct ttc cac gcc cgc ggt att ggc gtg atc atg gac tgg gtt cct 1123 Asp Ala Phe His Ala Arg Gly Ile Gly Val Ile Met Asp Trp Val Pro

335

1171

355

gcc cac ttc cct aag gat gat tgg gct ctt gcc cgc ttt gat ggc gaa

Ala His Phe Pro Lys Asp Asp Trp Ala Leu Ala Arg Phe Asp Gly Glu

350

330

345

					cct Pro											1219
					gac Asp											1267
					tac Tyr 395											1315
					gcc Ala											1363
					cca Pro											1411
					cag Gln											1459
					atc Ile											1507
					gac Asp 475											1555
					gac Asp											1603
cac His	cgc Arg	gca Ala	ttc Phe 505	cac His	cac His	agt Ser	gag Glu	ctc Leu 510	act Thr	ttc Phe	tcc Ser	ttg Leu	gtg Val 515	tac Tyr	gca Ala	1651
					gta Val											1699
					tgg Trp											1747
gcc Ala 550	gct Ala	ggt Gly	ctt Leu	cgc Arg	acc Thr 555	ttc Phe	ctt Leu	gcg Ala	tac Tyr	atg Met 560	tgg Trp	tca Ser	cac His	cca Pro	ggc Gly 565	1795
aag Lys	aag Lys	ctg Leu	ctt Leu	ttc Phe 570	atg Met	ggt Gly	cag Gln	gag Glu	ttt Phe 575	ggt Gly	cag Gln	cgt Arg	gaa Glu	gag Glu 580	tgg Trp	1843
gct Ala	gaa Glu	ggc Gly	cag Gln 585	gga Gly	ctg Leu	cca Pro	tgg Trp	gat Asp 590	att Ile	gtc Val	gac Asp	ggc Gly	tgg Trp 595	caa Gln	ggc Gly	1891

				cgc Arg									1939
				ctg Leu 620									1987
		_		gac Asp	-	-					_		2035
				ggc Gly									2083
				gag Glu									2131
				aac Asn									2179 .
				tcc Ser 700									2227
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<212> PRT

<213> Corynebacterium glutamicum

<400> 362

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Trp His Glu Thr Glu Ala Gly Ser Val Ile Arg Thr Arg Gln Val Gly 35 40 45

Ala Thr Gln Val Asn Leu Leu Ile Asp Asp Thr Ser His Val Met Thr 50 60

Pro Ile Gly Asp Asp Ile Phe Ala Ile Asp Leu Gly His Arg Glu Arg
65 70 75 80

Ala Asp Tyr Arg Leu Glu Val Thr Trp Pro Asp Gln Glu Pro Gln Val 85 90 95

Lys Ala Asp Pro Tyr Tyr Phe Leu Pro Thr Val Gly Glu Met Asp Ile 100 Tyr Leu Phe Ser Glu Gly Arg His Glu Arg Leu Trp Glu Ile Leu Gly Ala Asn Ile Lys Thr Tyr Gln Thr Ala Leu Gly Thr Val Arg Gly Thr 135 Ala Phe Thr Val Trp Ala Pro Asn Ala Ile Gly Cys Ala Val Val Gly 150 Gly Phe Asn Gly Trp Asn Ala Ser Gln His Pro Met Arg Ser Met Gly Gly Ser Gly Leu Trp Glu Leu Phe Ile Pro Gly Ile Glu Glu Gly Glu 185 Val Tyr Lys Phe Ala Val Gln Thr Arg Glu Gly Gln Arg Arg Asp Lys Ala Asp Pro Met Ala Arg Arg Ala Glu Leu Ala Pro Ala Thr Gly Ser 215 Ile Val Ala Ser Ser Glu Tyr Gln Trp Gln Asp Ser Glu Trp Leu Arg 235 Glu Arg Ser Gln Thr Asp Leu Ala Ser Lys Pro Met Ser Val Tyr Glu Val His Leu Gly Ser Trp Arg Trp Gly Lys Asn Tyr Glu Asp Leu Ala 265 Thr Glu Leu Val Asp Tyr Val Ala Asp Leu Gly Tyr Thr His Val Glu 280 Phe Leu Pro Val Ala Glu His Pro Phe Gly Gly Ser Trp Gly Tyr Gln 290 295 Val Thr Gly Tyr Tyr Ala Pro Thr Ser Arg Trp Gly Thr Pro Asp Gln 315 Phe Arg Ala Leu Val Asp Ala Phe His Ala Arg Gly Ile Gly Val Ile 325 Met Asp Trp Val Pro Ala His Phe Pro Lys Asp Asp Trp Ala Leu Ala Arg Phe Asp Gly Glu Ala Leu Tyr Glu His Pro Asp Trp Arg Arg Gly 365 Glu Gln Lys Asp Trp Gly Thr Leu Val Phe Asp Phe Gly Arg Asn Glu Val Arg Asn Phe Leu Val Ala Asn Ala Leu Tyr Trp Ile Glu Glu Phe 385 His Ile Asp Gly Leu Arg Val Asp Ala Val Ala Ser Met Leu Tyr Leu Asp Tyr Ser Arg Glu His Gly Glu Trp Glu Pro Asn Ile Tyr Gly Gly

 Arg Glu Asn Leu Glu Ala Val Gln Phe Leu Gln Glu Met Asn Ala Thr 435

 Val Leu Arg Leu His Pro Gly Asn Asn Thr 455

 Ser Trp Pro Gly Val Thr Ala Pro Thr Trp Asn Asn Asn Asn Asn Asn Asn Asn Thr 480

 Ser Leu Lys Asn Pro Val His Arg Ala Phe His His Ser Glu Leu Thr Phe

Ser Lys Asn Pro Val His Arg Ala Phe His His Ser Glu Leu Thr Phe 500 505 510

Ser Leu Val Tyr Ala Phe Ser Glu Arg Phe Val Leu Pro Ile Ser His 515 520 525

Asp Glu Val Val His Gly Lys Gly Ser Leu Trp Asp Arg Met Pro Gly 530 535 540

Asp Thr Trp Asn Lys Ala Ala Gly Leu Arg Thr Phe Leu Ala Tyr Met 545 550 555 560

Trp Ser His Pro Gly Lys Lys Leu Leu Phe Met Gly Gln Glu Phe Gly 565 570 575

Gln Arg Glu Glu Trp Ala Glu Gly Gln Gly Leu Pro Trp Asp Ile Val 580 585 590

Asp Gly Trp Gln Gly Glu Tyr His Glu Ala Ile Arg Thr Leu Thr Arg 595 600 605

Ser Leu Asn Gly Val Tyr Ser Asp Ser Pro Ala Leu His Thr Gln Asp 610 620

Phe Thr Gly Glu Gly Phe Thr Trp Asn Lys Gly Asp Asp Ala Thr Asn 625 630 635 640

Asn Ile Leu Ala Phe Thr Arg Phe Gly Ser Asp Gly Ser Gln Met Leu 645 650 655

Cys Val Phe Asn Leu Ser Gly Thr Ser Gln Pro Glu Tyr Gln Leu Gly 660 665 670

Val Ala Ala Gly Gly Glu Trp Lys Leu Val Leu Asn Thr Asp Asp Ala 675 680 685

Glu Phe Leu Gly Ala Glu Asn Asp Ile Ala Thr Ser Val Gln Ala Ala 690 695 700

Ala Thr Pro Arg Asp Asn Phe Ala Tyr Ser Leu Ser Leu His Val Pro 705 710 715 720

Ala Met Ser Ala Gln Phe Tyr Ser Leu Gln Lys

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ggc ttc ttt Gly Phe Phe 35	Ala Pro Ty						
ggc gcc gtg Gly Ala Val 50							
ggc ctc gaa Gly Leu Glu 65	Val Ile Le	g gat gtg u Asp Val 0	gtc tac Val Tyr	aac cac Asn His 75	acc gcc Thr Ala	Glu G	gc 240 ly 30
aac cac atg Asn His Met							
tac tac cga Tyr Tyr Arg							
ggt act ggt Gly Thr Gly 115	Asn Ser Le						
att atg gat Ile Met Asp 130							
ttc cgc ttc Phe Arg Phe 145		a Ser Thr					al
gac cgc ctg Asp Arg Leu							
tcc cag gtc Ser Gln Val							
tac caa gtg Tyr Gln Val 195	ggt aac tt Gly Asn Ph	c cca cca e Pro Pro 200	ctg tgg Leu Trp	Thr Glu	tgg aac Trp Asn 205	ggt aa Gly Ly	aa 624 7s

	_	_		-	_						gag Glu 220				672
											gat Asp				720
											gtg Val				768
					_	_	-	_			gag Glu	-		-	816
-				-			_		_		cac His		_		864
											gag Glu 300				912
-	_	_		-	-						ttg Leu	_	_		960
			_	_				-	-	_	gcc Ala	_			1008.
											ctg Leu				1056
	_	_	-	_	_		-	-	-		agc Ser			 -	1104
											agg Arg 380				1152
											gac Asp				1200
											gat Asp				1248
											ggc Gly				1296
											gac Asp				1344

atg ttc aac gct c Met Phe Asn Ala H 450					1392
cat ttc ggt atg a His Phe Gly Met L 465					1440
ggc cac ccg ctg g Gly His Pro Leu G 4					1488
gtt cct gcc cgt t Val Pro Ala Arg S 500			Gln Val Glu A		1536
tac acc aag ctt g Tyr Thr Lys Leu G 515	lu Glu Lys				1584
ctt gcg gca gag a Leu Ala Ala Glu L 530					1632
gca gca aag gaa g Ala Ala Lys Glu A 545					1680
gaa cgt gct tcg a Glu Arg Ala Ser T 5					1728
gat gcg att gcc g Asp Ala Ile Ala A 580				eu Pro Gln	1776
gat gaa gta gcg g Asp Glu Val Ala A 595	la Glu Val				1824
act gaa tot gac to Thr Glu Ser Asp So 610					1872
gcg gac gaa gaa g Ala Asp Glu Glu G 625		acaccg aaagt	ggcgt cgc		1913
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Arg Leu Arg Asp Le	eu Gly Met i	Arg Asn Tyr 25		n Ser Phe	

Gly Phe Phe Ala Pro Tyr Asn Asp Tyr Ala Ala Asn Lys Asn Pro Gly Gly Ala Val Ala Glu Phe Lys Gly Leu Val Arg Ser Tyr His Glu Ala Gly Leu Glu Val Ile Leu Asp Val Val Tyr Asn His Thr Ala Glu Gly Asn His Met Gly Pro Thr Ile Ala Phe Arg Gly Ile Asp Asn Glu Ala Tyr Tyr Arg Leu Val Glu Gly Asp Arg Arg His Tyr Met Asp Tyr Thr 105 Gly Thr Gly Asn Ser Leu Asn Val Arg Asp Pro His Ser Leu Gln Leu Ile Met Asp Ser Leu Arg Tyr Trp Val Thr Glu Met His Val Asp Gly Phe Arg Phe Asp Leu Ala Ser Thr Leu Ala Arg Glu Phe Asp Asp Val 155 Asp Arg Leu Ala Thr Phe Phe Asp Leu Val Gln Gln Asp Pro Val Val 165 170 Ser Gln Val Lys Leu Ile Ala Glu Pro Trp Asp Val Gly Glu Gly Gly 180 185 Tyr Gln Val Gly Asn Phe Pro Pro Leu Trp Thr Glu Trp Asn Gly Lys Tyr Arg Asp Thr Val Arg Asp Phe Trp Arg Gly Glu Pro Ala Thr Leu 215 Gly Glu Phe Ala Ser Arg Leu Thr Gly Ser Ser Asp Leu Tyr Ala Asn Asn Gly Arg Arg Pro Thr Ala Ser Ile Asn Phe Val Thr Ala His Asp 245 250 Gly Phe Thr Leu Asn Asp Leu Val Ser Tyr Asn Glu Lys His Asn Met 265 260 Ala Asn Gly Glu Asp Gly Arg Asp Gly Glu Ser His Asn Arg Ser Trp Asn Cys Gly Val Glu Gly Pro Thr Asp Asp Pro Glu Ile Met Gln Leu 290 Arg Ala Gln Gln Arg Arg Asn Phe Leu Thr Thr Leu Leu Leu Ser Gln 310 315 Gly Thr Pro Met Leu Ser His Gly Asp Glu Met Ala Arg Thr Gln Asn Gly Asn Asn Asn Val Tyr Cys Gln Asp Asn Glu Leu Ala Trp Val Asn 345

Trp Asp Gln Ala Glu Glu Asn Ala Asp Leu Val Ser Phe Thr Arg Arg

355 360 365

Leu Leu Arg Ile Arg Ala Asn His Pro Val Phe Arg Arg Gln Phe 370 375 380

Leu Ala Gly Gly Pro Leu Gly Ala Asp Val Arg Asp Arg Asp Ile Ala 385 390 395 400

Trp Leu Val Pro Asn Gly Thr Leu Met Thr Gln Asp Asp Trp Asp Phe 405 410 415

Ala Phe Gly Lys Ser Leu Gln Val Phe Phe Asn Gly Asp Ala Ile Glu 420 425 430

Glu Pro Asp Tyr Arg Gly Gln Lys Ile His Asp Asp Ser Phe Ile Leu 435 440 445

Met Phe Asn Ala His Phe Glu Pro Ile Asp Phe Asn Leu Pro Pro Glu 450 455 460

His Phe Gly Met Lys Trp Lys Leu Leu Val Asp Thr Thr Glu Ala Val 465 470 475 480

Gly His Pro Leu Glu Asp Leu Thr Ile Glu Ala Gly Gly Thr Ile Thr 485 490 495

Val Pro Ala Arg Ser Thr Met Leu Leu Arg Gln Val Glu Ala Pro Asp 500 505 510

Tyr Thr Lys Leu Glu Glu Lys Ile Ala Ala Glu Lys Arg Glu Gln Glu 515 520 525

Leu Ala Ala Glu Lys Glu Ala Ala Glu Lys Arg Glu Leu Glu Leu Ala 530 540

Ala Ala Lys Glu Ala Glu Asp Ala Ala Glu Ala Leu His Leu Ala Ala 545 550 555 560

Glu Arg Ala Ser Thr Gln Glu Ala Glu Leu Ala His Gln His Gly Ala 565 570 575

Asp Ala Ile Ala Asp Glu Val Ala Glu Glu Pro Gln Glu Leu Pro Gln 580 585 590

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Thr Glu Ser Asp Ser Glu Gln Ala Glu Val Ala Ser Glu Glu Pro Glu 610 615 620

Ala Asp Glu Glu Glu Lys 625 630

<210> 365

<211> 1496

<212> DNA

<213> Corynebacterium glutamicum

<220>

<221> CDS

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Ser 225	Phe	Thr	Arg	Arg	Leu 230	Leu	Arg	Ile	Arg	Ala 235	Asn	His	Pro	Val	Phe 240	
	cgc Arg															768
	cgc Arg															816
	gac Asp															864
	gat Asp 290															912
	tcc Ser															960
	ctc Leu															1008
	acc Thr															1056
	gga Gly															1104
	gag Glu 370															1152
	cgt Arg															1200
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	caa Gln															1344
	gag Glu 450															1392
	acc Thr															1440

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Val Gly Glu Gly Tyr Gln Val Gly Asn Phe Pro Pro Leu Trp Thr 50 55 60

Glu Trp Asn Gly Lys Tyr Arg Asp Thr Val Arg Asp Phe Trp Arg Gly
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Glu Pro Ala Thr Leu Gly Glu Phe Ala Ser Arg Leu Thr Gly Ser Ser 85 90 95

Asp Leu Tyr Ala Asn Asn Gly Arg Arg Pro Thr Ala Ser Ile Asn Phe
100 105 110

Val Thr Ala His Asp Gly Phe Thr Leu Asn Asp Leu Val Ser Tyr Asn 115 120 125

Glu Lys His Asn Met Ala Asn Gly Glu Asp Gly Arg Asp Gly Glu Ser 130 135 140

His Asn Arg Ser Trp Asn Cys Gly Val Glu Gly Pro Thr Asp Asp Pro 145 150 155 160

Glu Ile Met Gln Leu Arg Ala Gln Gln Arg Arg Asn Phe Leu Thr Thr 165 170 175

Leu Leu Ser Gln Gly Thr Pro Met Leu Ser His Gly Asp Glu Met
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Ala Arg Thr Gln Asn Gly Asn Asn Asn Val Tyr Cys Gln Asp Asn Glu 195 200 205

Leu Ala Trp Val Asn Trp Asp Gln Ala Glu Glu Asn Ala Asp Leu Val 210 215 220

Ser Phe Thr Arg Arg Leu Leu Arg Ile Arg Ala Asn His Pro Val Phe 225 230 235 240

Arg Arg Arg Gln Phe Leu Ala Gly Gly Pro Leu Gly Ala Asp Val Arg

245 250 255

Asp Arg Asp Ile Ala Trp Leu Val Pro Asn Gly Thr Leu Met Thr Gln 260 265 270

Asp Asp Trp Asp Phe Ala Phe Gly Lys Ser Leu Gln Val Phe Phe Asn 275 280 285

Gly Asp Ala Ile Glu Glu Pro Asp Tyr Arg Gly Gln Lys Ile His Asp 290 295 300

Asp Ser Phe Ile Leu Met Phe Asn Ala His Phe Glu Pro Ile Asp Phe 305 310 315 320

Asn Leu Pro Pro Glu His Phe Gly Met Lys Trp Lys Leu Leu Val Asp 325 330 335

Thr Thr Glu Ala Val Gly His Pro Leu Glu Asp Leu Thr Ile Glu Ala 340 345 350

Gly Gly Thr Ile Thr Val Pro Ala Arg Ser Thr Met Leu Leu Arg Gln 355 360 365

Val Glu Ala Pro Asp Tyr Thr Lys Leu Glu Glu Lys Ile Ala Ala Glu 370 375 380

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410
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His Gln His Gly Ala Asp Ala Ile Ala Asp Glu Val Ala Glu Glu Pro 435 440 445

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His Cys Tyr Leu Pro Gly Val Gln Pro Gly Gln Arg Tyr Gly Phe Arg 65 70 75 80	

Val His Gly Pro Trp Asn Pro Asp Glu Gly Lys Arg Cys Asp Ala Asn Lys Leu Leu Val Asp Pro Tyr Ala Arg Ala Phe Asp Gly Asp Phe Asp 105 Gly His Pro Ser Leu Phe Ser Tyr Asp Ile Thr Asn Pro Asn Asp Pro 115 Asn Gly Arg Asn Thr Glu Asp Ser Ile Asp His Thr Met Lys Ser Val 135 Val Val Asn Pro Phe 145 <210> 369 <211> 1635 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(1612) <223> RXN01550 <400> 369 ttcgccagca gtacttcttc acctctgctt ccctgcaggc catgattcag ggccacctgg 60 egeaceacaa ggaceteage aactttgeeg agtteactee gtg eag ete aat gae Val Gln Leu Asn Asp act cac cca gtg ttg gct atc cct gag ctt atg cgt ctg ctc atg gac 163 Thr His Pro Val Leu Ala Ile Pro Glu Leu Met Arg Leu Leu Met Asp 15 gag cat gac atg ggc tgg gaa gaa tcc tgg gca atc gtg ttc aag acc 211 Glu His Asp Met Gly Trp Glu Glu Ser Trp Ala Ile Val Phe Lys Thr 259 tte gea tae ace aac cae ace gtg etc ace gaa get ett gag eag tgg Phe Ala Tyr Thr Asn His Thr Val Leu Thr Glu Ala Leu Glu Gln Trp 45 gat cag cag atc ttc caa cag ctg ttc tgg cgc gtg tgg gaa atc atc 307 Asp Gln Gln Ile Phe Gln Gln Leu Phe Trp Arg Val Trp Glu Ile Ile aca gag atc gat cgc cgc ttc cgt ttg gag cgc gca gcc gat gga ctg 355 Thr Glu Ile Asp Arg Arg Phe Arg Leu Glu Arg Ala Ala Asp Gly Leu 70 . gat gaa gag acc atc gac cgc atg gct cca atc cag cac ggc act gtt 403 Asp Glu Glu Thr Ile Asp Arg Met Ala Pro Ile Gln His Gly Thr Val cat atg gca tgg att gcc tgt tac gcg gca tat tcc atc aat ggc gtg 451 His Met Ala Trp Ile Ala Cys Tyr Ala Ala Tyr Ser Ile Asn Gly Val 105 110 115

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Ala Leu Glu Gln Trp Asp Gln Gln Ile Phe Gln Gln Leu Phe Trp Arg 50 55 60

Val Trp Glu Ile Ile Thr Glu Ile Asp Arg Phe Arg Leu Glu Arg 65 70 75 80

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Lys Arg Ala Leu Asp Ala Leu Asp Asn Gly Thr Leu Asn Asp Asn Asn Ser Gly Leu Phe Tyr Asp Leu Lys His Ser Leu Ile His Gly Tyr Gly 420 425 Lys Asp Ala Ser Asp Thr Tyr Tyr Val Leu Gly Asp Phe Ala Asp Tyr 440 Arg Glu Thr Arg Asp Arg Met Ala Ala Asp Tyr Ala Ser Asp Pro Leu 455 Gly Trp Ala Arg Met Ala Trp Ile Asn Ile Cys Glu Ser Gly Arg Phe 470 Ser Ser Asp Arg Thr Ile Arg Asp Tyr Ala Thr Glu Ile Trp Lys Leu 490 Glu Pro Thr Pro Ala Val Lys Lys 500 <210> 371 <211> 1367 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (1)..(1344) <223> FRXA01550 <400> 371 ato tto caa cag ctg tto tgg cgc gtg tgg gaa ato ato aca gag ato 48 Ile Phe Gln Gln Leu Phe Trp Arg Val Trp Glu Ile Ile Thr Glu Ile gat cgc cgc ttc cgt ttg gag cgc gca gcc gat gga ctg gat gaa gag 96 Asp Arg Arg Phe Arg Leu Glu Arg Ala Ala Asp Gly Leu Asp Glu Glu 20 acc atc gac egc atg get eca atc eag ege gge act gtt eat atg gea 144 Thr Ile Asp Arg Met Ala Pro Ile Gln Arg Gly Thr Val His Met Ala 35 tgg att gcc tgt tac gcg gca tat tcc atc aat ggc gtg gca gcg ctg 192 Trp Ile Ala Cys Tyr Ala Ala Tyr Ser Ile Asn Gly Val Ala Ala Leu 50 cac acc gag atc atc aag gcc gag acc ttg gct gac tgg tac gca ctg His Thr Glu Ile Ile Lys Ala Glu Thr Leu Ala Asp Trp Tyr Ala Leu 65 70 tgg cca gag aag ttc aac aac aag act aac ggt gtt acc cca cgc cgt 288 Trp Pro Glu Lys Phe Asn Asn Lys Thr Asn Gly Val Thr Pro Arg Arg 90 95 tgg ctg cgc atg atc aac cca ggt ctg tct gac ctg ctc act cga ctt 336 Trp Leu Arg Met Ile Asn Pro Gly Leu Ser Asp Leu Leu Thr Arg Leu 100 105

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Gly Ile Glu Ile Asp Pro Glu Ser Ile Phe Asp Val Gln Ile Lys Arg Leu His Glu Tyr Lys Arg Gln Leu Met Asn Ala Leu Tyr Val Leu Asp 185 Leu Tyr Phe Arg Ile Lys Glu Asp Gly Leu Thr Asp Ile Pro Ala Arg Thr Val Ile Phe Gly Ala Lys Ala Ala Pro Gly Tyr Val Arg Ala Lys Ala Ile Ile Lys Leu Ile Asn Ser Ile Ala Asp Leu Val Asn Asn Asp Pro Glu Val Ser Pro Leu Leu Lys Val Val Phe Val Glu Asn Tyr Asn 250 Val Ser Pro Ala Glu His Ile Leu Pro Ala Ser Asp Val Ser Glu Gln 265 260 Ile Ser Thr Ala Gly Lys Glu Ala Ser Gly Thr Ser Asn Met Lys Phe 280 Met Met Asn Gly Ala Leu Thr Leu Gly Thr Met Asp Gly Ala Asn Val 295 Glu Ile Val Asp Ser Val Gly Glu Glu Asn Ala Tyr Ile Phe Gly Ala 310 315 Arg Val Glu Glu Leu Pro Ala Leu Arg Glu Ser Tyr Glu Pro Tyr Glu Leu Tyr Glu Thr Val Pro Gly Leu Lys Arg Ala Leu Asp Ala Leu Asp Asn Gly Thr Leu Asn Asp Asn Asn Ser Gly Leu Phe Tyr Asp Leu Lys 360 His Ser Leu Ile His Gly Tyr Gly Lys Asp Ala Ser Asp Thr Tyr Tyr 370 Val Leu Gly Asp Phe Ala Asp Tyr Arg Glu Thr Arg Asp Arg Met Ala 395 Ala Asp Tyr Ala Ser Asp Pro Leu Gly Trp Ala Arg Met Ala Trp Ile 405

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170

Leu Arg Asn Val Thr Asp Arg Leu Tyr Gly Gly Asp Asn Glu His Arg

155

145

Ile Lys Gln Glu Leu Val Leu Gly Val Gly Val Arg Ala Val Asn 180 185 Ala Phe Cys Glu Ala Arg Gly Leu Lys Arg Ser Ser Val Ala His Leu Asn Glu Gly His Ala Gly Phe Leu Thr Leu Glu Arg Ile Arg Glu Arg Ile Ala Glu Gly Met Glu Tyr Pro Ala Ala Phe Glu Gln Val Arg Ala Ser Asn Ile Phe Thr Thr His Thr Pro Val Pro Ala Gly Ile Asp Arg Phe Asp Met Glu Met Val Arg Arg Tyr Leu Gly Gly Gln Pro Glu Asp Gln Gln Leu Cys Val Gly Val Pro Ile Glu Lys Ala Leu Glu Leu Gly Gln Glu Ser Asp Pro His Arg Phe Asn Met Ala His Met Gly Leu Arg Ala Ser Gln His Ala Asn Gly Val Ala Lys Leu His Gly Glu Val 310 315 Ser Arg Asp Met Phe Ala Gly Leu Tyr Pro Gly Tyr Glu Pro Arg Glu Val Pro Ile Gly His Val Thr Asn Gly Val His Leu Pro Thr Trp Val 345 Lys Pro Glu Met Lys Glu Leu Ile Asp Arg Val Thr Gly Gly Ala Asp 355 360 Leu Ala Val Ala Asp Ser Trp Ser Asn Pro Gln Ala Val Glu Ser Glu 375 Lys Ile Trp Lys Val Arg Asn Lys Phe Arg Ala Asp Leu Val Glu Val 395 Ala Arg Ala Ala Thr Ala Lys Ser Trp Ser His Arg Gly His Thr Glu 405 Ala Glu Leu Ala Trp Thr Ser Arg Val Leu Asp Pro Asn Val Leu Thr 425 Ile Gly Phe Ala Arg Arg Val Ser Thr Tyr Lys Arg Leu Thr Leu Met Leu Arg Asn Pro Glu Arg Leu Arg Ser Ile Leu Leu Asn Glu Glu Arg 450 Pro Val Gln Phe Val Ile Ala Gly Lys Ala His Pro His Asp Met Gly Gly Lys Lys Leu Met Gln Glu Ile Val His Phe Ala Asp Gln Ala Gly

490

Val Arg Asp Arg Phe Leu Phe Leu Pro Asp Tyr Asp Ile Asn Leu Ala 500 505 510

Ser Tyr Leu Ile Ser Gly Ala Asp Val Trp Leu Asn Asn Pro Val Arg 515 520 525

Pro Gln Glu Ala Ser Gly Thr Ser Gly Met Lys Ala Val Met Asn Gly 530 535 540

Gly Leu Thr Leu Ser Ile Ser Asp Gly Trp Trp Asp Glu Met Pro Lys 545 550 560

Glu Thr Thr Gly Trp Thr Ile Pro Thr Val Glu Ser Gln Asp Leu Glu 565 570 575

Cys Arg Asp His Leu Glu Ser Gln Ala Leu Tyr Asp Leu Leu Glu Asn 580 585 590

Glu Val Ala Pro Leu Phe Tyr Lys Arg Asp Lys Asn Gly Ile Pro Gln 595 600 605

Asp Trp Leu Asp Leu Val Arg Glu Ser Trp Thr Thr Leu Ser Pro Met 610 620

Val Thr Ser Thr Arg Met Val Arg Asp Tyr Thr Thr Gln Tyr Tyr Arg 625 630 635 640

Pro Thr Lys His Gln Ala Glu Leu Ile Ala Gln Pro Ala Glu Ala Ala 645 650 655

Asp Tyr Ala Ala Trp Leu Glu His Ile Lys Ala Glu Trp Ala Gly Val 660 665 670

Lys Val Ser Asp Leu Lys Ile Ser Glu Ser Ala Ile Thr Ala Gln Glu 675 680 685

Leu Glu Val Ser Val Arg Val Asp Ser Gly Ser Leu Asn Asp Asp Glu 690 695 700

Phe Gln Ala Gln Ala Leu Phe Gly Ala Leu Gly His Asn Gly Asp Ile 705 710 715 720

Glu Asp Pro Glu Ile Thr Val Leu Thr Pro Arg Gly Asp Gly Ala Tyr 725 730 735

Ala Ala Lys Val Ser Thr Asp Leu Pro Gly Asn Tyr Gly Ile Thr Ala 740 745 750

Arg Val Val Pro Asn Asn Arg Met Leu Val Ser Pro Ala Glu Thr Arg 755 760 765

Leu Ile Thr Tyr Leu Glu Asn
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<213> Corynebacterium glutamicum

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cgc gtt gat Arg Val Asp 225											720
ctc ttt ggt Leu Phe Gly											768
acc gtt ttg Thr Val Leu				y Ala							816
act gac ctg Thr Asp Leu 275	Pro Gly										864
aac agg atg Asn Arg Met 290											912
gag aac tag Glu Asn 305	ggcgaaa d	ctagcttt	ac caa								941
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Ile Ala Gly	5		Asp Gl	10	í				15		
Ile Ala Gly 1	Val His	Phe Ala	Asp Gl 2	10 n Ala 5	Gly	Val	Arg	Asp 30	15 Arg	Phe	
Ile Ala Gly 1 Gln Glu Ile Leu Phe Leu	Val His 20 Pro Asp	Phe Ala	Asp Gl 2 Ile As 40	10 n Ala 5 n Leu	Gly	Val Ser	Arg Tyr 45	Asp 30 Leu	15 Arg Ile	Phe Ser	
Ile Ala Gly 1 Gln Glu Ile Leu Phe Leu 35 Gly Ala Asp	Val His 20 Pro Asp Val Trp	Phe Ala Tyr Asp Leu Asn 55	Asp Gl 2 Ile As 40 Asn Pr	10 n Ala 5 n Leu o Val	Gly Ala Arg	Val Ser Pro 60	Arg Tyr 45 Gln	Asp 30 Leu Glu	15 Arg Ile Ala	Phe Ser Ser	
Ile Ala Gly 1 Gln Glu Ile Leu Phe Leu 35 Gly Ala Asp 50 Gly Thr Ser	Val His 20 Pro Asp Val Trp Gly Met	Phe Ala Tyr Asp Leu Asn 55 Lys Ala 70	Asp Gl 2 Ile As 40 Asn Pr	n Ala 5 n Leu o Val	Gly Ala Arg Gly 75	Val Ser Pro 60 Gly	Arg Tyr 45 Gln Leu	Asp 30 Leu Glu Thr	15 Arg Ile Ala Leu	Phe Ser Ser Ser 80	
Ile Ala Gly 1 Gln Glu Ile Leu Phe Leu 35 Gly Ala Asp 50 Gly Thr Ser 65	Val His 20 Pro Asp Val Trp Gly Met Gly Trp 85	Phe Ala Tyr Asp Leu Asn 55 Lys Ala 70 Trp Asp	Asp Gl 2 Ile As 40 Asn Pr Val Me	n Ala 5 n Leu O Val t Asn t Pro 90 p Leu	Gly Ala Arg Gly 75 Lys	Val Ser Pro 60 Gly Glu	Arg Tyr 45 Gln Leu Thr	Asp 30 Leu Glu Thr	15 Arg Ile Ala Leu Gly 95	Phe Ser Ser Ser Trp	
Ile Ala Gly 1 Gln Glu Ile Leu Phe Leu 35 Gly Ala Asp 50 Gly Thr Ser 65 Ile Ser Asp	Val His 20 Pro Asp Val Trp Gly Met Gly Trp 85 Thr Val	Phe Ala Tyr Asp Leu Asn 55 Lys Ala 70 Trp Asp Glu Ser	Asp Gl 2 Ile As 40 Asn Pr Val Me Glu Me	n Ala 5 n Leu O Val t Asn t Pro 90 p Leu 5	Gly Ala Arg Gly 75 Lys	Val Ser Pro 60 Gly Glu Cys	Arg Tyr 45 Gln Leu Thr	Asp 30 Leu Glu Thr Thr	15 Arg Ile Ala Leu Gly 95 His	Phe Ser Ser 80 Trp Leu	
Ile Ala Gly 1 Gln Glu Ile Leu Phe Leu 35 Gly Ala Asp 50 Gly Thr Ser 65 Ile Ser Asp Thr Ile Pro Glu Ser Gln	Val His 20 Pro Asp Val Trp Gly Met Gly Trp 85 Thr Val 100 Ala Leu	Phe Ala Tyr Asp Leu Asn 55 Lys Ala 70 Trp Asp Glu Ser Tyr Asp	Asp Gl 2 Ile As 40 Asn Pr Val Me Glu Me Glu As 10 Leu Le 120	n Ala 5 n Leu O Val t Asn t Pro 90 p Leu 5	Gly Ala Arg Gly 75 Lys Glu Asn	Val Ser Pro 60 Gly Glu Cys	Arg Tyr 45 Gln Leu Thr Arg Val 125	Asp 30 Leu Glu Thr Thr Asp 110 Ala	15 Arg Ile Ala Leu Gly 95 His	Phe Ser Ser 80 Trp Leu Leu	

Met Val Arg Asp Tyr Thr Thr Gln Tyr Tyr Arg Pro Thr Lys His Gln 165 170 Ala Glu Leu Ile Ala Gln Pro Ala Glu Ala Asp Tyr Ala Ala Trp 180 Leu Glu His Ile Lys Ala Glu Trp Ala Gly Val Lys Val Ser Asp Leu 200 Lys Ile Ser Glu Ser Ala Ile Thr Ala Gln Glu Leu Glu Val Ser Val Arg Val Asp Ser Gly Ser Leu Asn Asp Asp Glu Phe Gln Ala Gln Ala 230 235 Leu Phe Gly Ala Leu Gly His Asn Gly Asp Ile Glu Asp Pro Glu Ile 245 250 Thr Val Leu Thr Pro Arg Gly Asp Gly Ala Tyr Ala Ala Lys Val Ser Thr Asp Leu Pro Gly Asn Tyr Gly Ile Thr Ala Arg Val Val Pro Asn 280 Asn Arg Met Leu Val Ser Pro Ala Glu Thr Arg Leu Ile Thr Tyr Leu 290 295 Glu Asn 305 <210> 377 <211> 1206 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (1)..(1206) <223> FRXA02113 <400> 377 cta ggt cga atc aac gcc gag gag caa aac ctc agc gaa tac ctc agc 48 Leu Gly Arg Ile Asn Ala Glu Glu Gln Asn Leu Ser Glu Tyr Leu Ser gac aag ctg tgg tac cag gac acc gca gat gca acc gat gct gtc gga 96 Asp Lys Leu Trp Tyr Gln Asp Thr Ala Asp Ala Thr Asp Ala Val Gly gat cca ctc gtt gcg tac ttc tcc atg gag ttt ggc att cac cca agc 144 Asp Pro Leu Val Ala Tyr Phe Ser Met Glu Phe Gly Ile His Pro Ser ctg cca atc tac tct ggc gga ctt ggt gtg ctt gcg ggc gag aac atg 192 Leu Pro Ile Tyr Ser Gly Gly Leu Gly Val Leu Ala Gly Glu Asn Met 50 55 aag tot goa tot gac ttg ggt gtg coa ctg atc ggt gtt ggt ttg ctc Lys Ser Ala Ser Asp Leu Gly Val Pro Leu Ile Gly Val Gly Leu Leu 65

			cag Gln						288
			gat Asp						336
			cag Gln 120						384
			gca Ala						432
			acc Thr						480
			ctg Leu						528
			ggt Gly						576
			ctg Leu 200						624
			ctg Leu						672
			cca Pro						720
			acc Thr	-	_		-	_	768
			cgt Arg						816
			gtt Val 280						864
			cgc Arg						912
			ggc Gly						960

	cgt Arg								1008
	ccc Pro								1056
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gct Ala				-					1206

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<400> 378

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Asp Pro Leu Val Ala Tyr Phe Ser Met Glu Phe Gly Ile His Pro Ser 35 40 45

Leu Pro Ile Tyr Ser Gly Gly Leu Gly Val Leu Ala Gly Glu Asn Met 50 55 60

Lys Ser Ala Ser Asp Leu Gly Val Pro Leu Ile Gly Val Gly Leu Leu 65 70 75 80

Tyr Thr His Gly Tyr Phe Thr Gln Ser Leu Ser Gly Asp Gly Trp Gln
85 90 95

Gln Glu Glu Tyr Lys Tyr His Asp Pro Ala Glu Leu Pro Ile Glu Ala 100 105 110

Val Lys Asp Lys Asn Gly Glu Gln Val Thr Val Ser Val Thr Tyr Pro 115 120 125

Gly Ala Gln Glu Val Lys Ile Ala Leu Trp Val Ala Asn Val Gly Arg 130 135 140

Ile Pro Leu Leu Leu Asp Thr Asn Ile Glu Ala Asn Pro Glu Glu 145 150 155 160

Leu Arg Asn Val Thr Asp Arg Leu Tyr Gly Gly Asp Asn Glu His Arg 165 170 175

Ile Lys Gln Glu Leu Val Leu Gly Val Gly Gly Val Arg Ala Val Asn 180 185 190

Ala Phe Cys Glu Ala Arg Gly Leu Lys Arg Ser Ser Val Ala His Leu 195 200 205

Asn Glu Gly His Ala Gly Phe Leu Thr Leu Glu Arg Ile Arg Glu Arg 210 215 220

Ile Ala Glu Gly Met Glu Tyr Pro Ala Ala Phe Glu Gln Val Arg Ala 225 230 235 240

Ser Asn Ile Phe Thr Thr His Thr Pro Val Pro Ala Gly Ile Asp Arg 245 250 255

Phe Asp Met Glu Met Val Arg Arg Tyr Leu Gly Gly Gly Gln Pro Glu 260 265 270

Asp Gln Gln Leu Cys Val Gly Val Pro Ile Glu Lys Ala Leu Glu Leu 275 280 285

Gly Gln Glu Ser Asp Pro His Arg Phe Asn Met Ala His Met Gly Leu 290 295 300

Arg Ala Ser Gln His Ala Asn Gly Val Ala Lys Leu His Gly Glu Val 305 310 315 320

Ser Arg Asp Met Phe Ala Gly Leu Tyr Pro Gly Tyr Glu Pro Arg Glu 325 330 335

Val Pro Ile Gly His Val Thr Asn Gly Val His Leu Pro Thr Trp Val 340 345 350

Lys Pro Glu Met Lys Glu Leu Ile Asp Arg Val Thr Gly Gly Ala Asp 355 360 365

Leu Ala Val Ala Asp Ser Trp Ser Asn Pro Gln Ala Val Glu Ser Glu 370 375 380

Lys Ile Trp Lys Val Arg Asn Lys Phe Arg Ala Asp Leu Val Glu Val 385 390 395 400

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tgg gt Trp Va															163
gtg co Val Pi															211
atc ga Ile Gl		Val													259
cag ct Gln Le															307
cag ga Gln Gl 70															355
aat gt Asn Va	c gaa al Glu	gct Ala	tat Tyr 90	cgt Arg	tcg Ser	gag Glu	atc Ile	aat Asn 95	cgg Arg	atc Ile	gct Ala	cag Gln	gcg Ala 100	aag Lys	.403
tat co															451
gat co Asp Pr		Asn													499
aag to Lys Se 13	er Thr														547
tcc go Ser Al 150															595
ttc ca Phe Gl															643
aag ga Lys Gl															691
gcc ct Ala Le															739
ttg ga Leu As 21	p Ile														787

ggt gcg gtg Gly Ala Val 230												835
ggt ggc att Gly Gly Ile												883
gcg tat cag Ala Tyr Gln												931
atg gct ggc Met Ala Gly 280	Gly Thr											979
gtc att gga Val Ile Gly 295			y Āla									1027
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155

Leu Ser Thr Leu Thr Lys Ser Thr Ser Asp Val Val Glu Ser Leu Asn 130 135 140

Ala Glu Thr Glu Lys Ser Ala Glu Ala Val Tyr Gln Ala Asn Arg Thr

Lys Ala Glu Ala Glu Phe Gln Leu Gly Gln Leu Lys Val Arg Gln Ala 165 170 175

150

Glu Leu Glu Ser Glu Lys Glu Ala Leu Asp Gly Arg Lys Ser Glu Ile 180 185 190

Arg Asp Arg Val Asp Ala Leu Thr Pro Gln Glu Arg Glu Met Trp Val 195 200 205

Ala Lys Asn Gly Pro Leu Asp Ile Asp Leu Thr Asp Leu Leu Gly Leu 210 215 220

Ser Ala Ala Thr Ser Gly Ala Val Asp Ala Ala Leu Ser Lys Leu Gly 225 230 235 240

Ser Pro Tyr Gly Trp Gly Gly Ile Gly Pro Asn Glu Phe Asp Cys Ser 245 250 255

Gly Leu Ile Tyr Trp Ala Tyr Gln Gln Met Gly Lys Thr Leu Pro Arg 260 265 270

Thr Ser Gln Ala Gln Met Ala Gly Gly Thr Pro Val Ser Arg Asp Glu 275 280 285

Leu Gln Pro Gly Asp Val Ile Gly Tyr Tyr Pro Gly Ala Thr His Val 290 295 300

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<223> RXA01478

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Met Thr Ile Pro Gly

1 5

get tee aca cag act gat ate eet etg gae aca ett ett gag gat tae 163

Ala	Ser	Thr	Gln	Thr 10	Asp	Ile	Pro	Leu	Asp 15	Thr	Leu	Leu	Glu	Asp 20	Tyr	
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					tgc Cys											259
					gat Asp											307
					atc Ile 75											355
					cgt Arg											403
					ggt Gly											451
					gtg Val											499
					tat Tyr											547
					agc Ser 155											595
					cct Pro											643
					cac His											691
					tcg Ser											739
					agc Ser											787
gag Glu 230	ctc Leu	ccc Pro	cac His	cag Gln	cgc Arg 235	ctc Leu	tac Tyr	gac Asp	gct Ala	gaa Glu 240	gtc Val	cgc Arg	cgc Arg	tcc Ser	atg Met 245	835
					ttg Leu											883

250 255 260 gca ccg acc acc tca cta cca gag gat ttc gga ggc atc cgt aac tgg Ala Pro Thr Thr Ser Leu Pro Glu Asp Phe Gly Gly Ile Arg Asn Trp 265 270 gac tac cgc tac gtg tgg ctg cgc gac tcc gca ctc acc att gaa gcc 979 Asp Tyr Arg Tyr Val Trp Leu Arg Asp Ser Ala Leu Thr Ile Glu Ala ctc gtg gaa tac gga ttc tcc caa gca gcc ctc caa tgg cgc acc tgg 1027 Leu Val Glu Tyr Gly Phe Ser Gln Ala Ala Leu Gln Trp Arg Thr Trp 295 300 ctg ctg cgc gcc atc gca ggc gac ccg gaa aac ctc cgc atc atg tat 1075 Leu Leu Arg Ala Ile Ala Gly Asp Pro Glu Asn Leu Arg Ile Met Tyr 310 315 ggc ctc ggc ggc gaa cga cac ctc cct gaa cgc gaa ctc caa cac ctg 1123 Gly Leu Gly Gly Glu Arg His Leu Pro Glu Arg Glu Leu Gln His Leu 330 335 cgc gga tac gaa aac tcc gtg cct gtt cgc gtt ggc aat gga gcc gcc 1171 Arg Gly Tyr Glu Asn Ser Val Pro Val Arg Val Gly Asn Gly Ala Ala 345 gaa caa tac caa gca gat gtc gtc ggc gaa gta atg gtc gcg ctt gaa 1219 Glu Gln Tyr Gln Ala Asp Val Val Gly Glu Val Met Val Ala Leu Glu 360 365 acc atc cgc cgc gcc ggg tgc ctc gag gac gaa ttc tcc tgg ggc atg 1267 Thr Ile Arg Arg Ala Gly Cys Leu Glu Asp Glu Phe Ser Trp Gly Met 375 380 caa aaa gcc atc ctc gat ttc caa gaa gcc aac ttc gac cgc aag gat 1315 Gln Lys Ala Ile Leu Asp Phe Gln Glu Ala Asn Phe Asp Arg Lys Asp 395 caa ggc atc tgg gaa atg cgc tcc gaa ccg caa tat ttc acc cac ggc 1363 Gln Gly Ile Trp Glu Met Arg Ser Glu Pro Gln Tyr Phe Thr His Gly 410 415 cgc gcc atg atg tgg gcc ggc ttc gac cgc ggc atc aaa gcc atc gaa 1411 Arg Ala Met Met Trp Ala Gly Phe Asp Arg Gly Ile Lys Ala Ile Glu 430 gaa ttc aac ctc gac ggc ccc atc gag cgc tgg cgt gaa ctc cgc gcc 1459 Glu Phe Asn Leu Asp Gly Pro Ile Glu Arg Trp Arg Glu Leu Arg Ala 445 aaa ctc cgc gaa gaa atc atg acc aac ggc ttc aac gaa gag atc caa 1507 Lys Leu Arg Glu Glu Ile Met Thr Asn Gly Phe Asn Glu Glu Ile Gln tcc ttc acc cag tgc tac gac acc caa gtc gac gcc tcg ctt 1555 Ser Phe Thr Gln Cys Tyr Asp Asn Thr Gln Val Asp Ala Ser Leu Leu 470 480 cag etc gec caa ata gge tte atc gge tte gac gat eca aaa atg etc 1603 Gln Leu Ala Gln Ile Gly Phe Ile Gly Phe Asp Asp Pro Lys Met Leu

495

490

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ctt cac agg t Leu His Arg T 520	ac cac a yr His T	acc gac Thr Asp	ggg tc Gly Se 525	gac Asp	ggc Gly	ctt Leu	gcc Ala 530	ggc Gly	gac Asp	gaa Glu	1699
tac ccc ttc c Tyr Pro Phe L 535											1747
tcc aac cgc c Ser Asn Arg L 550	eu Asp G										1795
gtc caa agc c Val Gln Ser P	ca ctt g ro Leu G 570	gc cta Sly Leu	ctg gc Leu Al	gag Glu 575	gaa Glu	tac Tyr	tcc Ser	acc Thr	cac His 580	cat His	1843
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atc agc gct g Ile Ser Ala A 600											1936
tagagtctaa gg	tgtcattc	ttg									1959
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Val	Glu 130	Ser	Ile	Leu	Arg	Leu 135	Arg	Phe	Asp	Tyr	Gly 140	Glu	Ser	Thr	Pro
Tyr 145	Phe	Arg	Thr	Ser	Thr 150	Val	Asp	Gly	Ile	Ser 155	Ile	Val	Gln	Ala	Val
Ala	Gly	Pro	Asn	Ala 165	Val	Tyr	Val	Arg	Gly 170	Pro	Glu	Met	Pro	His 175	Arç
Pro	Ala	Lys	Asp 180	Cys	His	Ser	Gly	Thr 185	Phe	His	Leu	Thr	Ala 190	Gly	Glu
Ser	Val	Glu 195	Trp	Val	Leu	Thr	Trp 200	Ala	Pro	Ser	Phe	Glu 205	Pro	His	Pro
Pro	Met 210	Pro	Asp	Tyr	Thr	Arg 215	Ser	Leu	Glu	Ser	Thr 220	Leu	Ser	Phe	Trp
Ala 225	Ser	Trp	Val	Glu	Glu 230	Leu	Pro	His	Gln	Arg 235	Leu	Tyr	Asp	Ala	Glu 240
Val	Arg	Arg	Ser	Met 245	Leu	Val	Leu	Arg	Ala 250	Leu	Thr	Asp	Leu	Gln 255	Thr
Gly	Gly	Ile	Val 260	Ala	Ala	Pro	Thr	Thr 265	Ser	Leu	Pro	Glu	Asp 270	Phe	Gly
Gly	Ile	Arg 275	Asn	Trp	Asp	Tyr	Arg 280	Tyr	Val	Trp	Leu	Arg 285	Asp	Ser	Ala
Leu	Thr 290	Ile	Glu	Ala	Leu	Val 295	Glu	Tyr	Gly	Phe	Ser 300	Gln	Ala	Ala	Leu
Gln 305	Trp	Arg	Thr	Trp	Leu 310	Leu	Arg	Ala	Ile	Ala 315	Gly	Asp	Pro	Glu	Asn 320
Leu	Arg	Ile	Met	Tyr 325	Gly	Leu	Gly	Gly	Glu 330	Arg	His	Leu	Pro	Glu 335	Arg
Glu	Leu	Gln	His 340	Leu	Arg	Gly	Tyr	Glu 345	Asn	Ser	Val	Pro	Val 350	Arg	Val
Gly	Asn	Gly 355	Ala	Ala	Glu	Gln	Tyr 360	Gln	Ala	Asp	Val	Val 365	Gly	Glu	Val
Met	Val 370	Ala	Leu	Glu	Thr	Ile 375	Arg	Arg	Ala	Gly	Cys 380	Leu	Glu	Asp	Glu
Phe 385	Ser	Trp	Gly	Met	Gln 390	Lys	Ala	Ile	Leu	Asp 395	Phe	Gln	Glu	Ala	Asn 400
Phe	Asp	Arg	Lys	Asp 405	Gln	Gly	Ile	Trp	Glu 410	Met	Arg	Ser	Glu	Pro 415	Gln
Tyr	Phe	Thr	His 420	Gly	Arg	Ala	Met	Met 425	Trp	Ala	Gly	Phe	Asp 430	Arg	Gly
Ile	Lys	Ala 435	Ile	Glu	Glu	Phe	Asn 440	Leu	Asp	Gly	Pro	Ile 445	Glu	Arg	Trp

Arg Glu Leu Arg Ala Lys Leu Arg Glu Glu Ile Met Thr Asn Gly Phe Asn Glu Glu Ile Gln Ser Phe Thr Gln Cys Tyr Asp Asn Thr Gln Val 470 Asp Ala Ser Leu Cln Leu Ala Gln Ile Gly Phe Ile Gly Phe Asp 485 Asp Pro Lys Met Leu Ser Thr Val Ala Arg Ile Glu Gln Glu Leu Leu 500 505 Asp Ala His Gly Phe Leu His Arg Tyr His Thr Asp Gly Ser Asp Gly 520 Leu Ala Gly Asp Glu Tyr Pro Phe Leu Ile Cys Ser Phe Trp Leu Val 535 530 Glu Gln Tyr Ala Ser Ser Asn Arg Leu Asp Glu Ala Lys Glu Lys Met 550 555 Asn Arg Ile Leu Ala Val Gln Ser Pro Leu Gly Leu Leu Ala Glu Glu Tyr Ser Thr His His Gly Arg Leu Ala Gly Asn Tyr Pro Gln Ala Phe 585 Ser His Ile Gly Leu Ile Ser Ala Ala Arg Ala Ile Asn Phe Glu Glu 595 600 Ala Arg Asn Arg 610 <210> 383 <211> 658 <212> DNA <213> Corynebacterium glutamicum <220> <221> CDS <222> (101)..(658) <223> RXA01888 <400> 383 agtagatact agataccacc cattgatgcc gtcaaggggt ttcctgtaaa gatgtaagag 60 attaagaaaa gaggtagata tggcgtcaaa gcgaccgaca atg gct gat gtg gca Met Ala Asp Val Ala aaa get get gga gta tee aet geg etg gte tee ate gtg ttt ege gat Lys Ala Ala Gly Val Ser Thr Ala Leu Val Ser Ile Val Phe Arg Asp 10 20 gcc ccc gga gca agt gaa tcc acc cgc aac cat gtg aaa gaa aaa gcc Ala Pro Gly Ala Ser Glu Ser Thr Arg Asn His Val Lys Glu Lys Ala 25 30 35 gcc gaa ctc gga tac att cct gat cga cga gcc caa aaa ctt cgc caa. Ala Glu Leu Gly Tyr Ile Pro Asp Arg Arg Ala Gln Lys Leu Arg Gln

40 . 45 50

							gtg Val					His				307
							ctc Leu									355
							atc Ile									403
							gaa Glu									451
							gat Asp 125									499
							ggt Gly									547
							ggc Gly		Gln							595
atc Ile	gaa Glu	tta Leu	ggc Gly	cac His 170	gaa Glu	cac His	atc Ile	atc Ile	tac Tyr 175	atc Ile	gat Asp	ggt Gly	ggc Gly	gac Asp 180	gcc Ala	643
	ggc Gly															658

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<213> Corynebacterium glutamicum

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Val Lys Glu Lys Ala Ala Glu Leu Gly Tyr Ile Pro Asp Arg Ala 35 40 45

Gln Lys Leu. Arg Gln Asn Arg Ser Gly Leu Ile Gly Val Ala Phe Glu 50 55 60

Met His Gln Ala Phe His Gly Asp Ile Val Glu His Leu Tyr Pro Thr 65 70 75 80

Ala Arg Lys His Gly Phe Asp Leu Tyr Leu Ser Ala Ile Thr Pro Thr

Arg Thr Glu Lys Asp, Ala Val Asn Ala Leu Ile Arg Glu Arg Cys Glu
100 105 110

Ala Val Ile Leu Leu Gly Ser Arg Met Ser Pro Ser Asp Leu Glu Thr 115 120 125

Ile Ala Gln Gln Leu Pro Val Gln Val Ile Ala Arg Gly Ser Gly Thr 130 135 140

Pro Lys Val Ser Ser Val His Val Asp Asp Ala Val Gly Ala Gln Leu 145 150 155 160

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Asp Gly Gly Asp Ala Pro Gly Thr Gln Glu 180 185

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gga atc gat agt tcc acc caa tcc tgc aag gct ttg ctt gtc gac gcc 163
Gly Ile Asp Ser Ser Thr Gln Ser Cys Lys Ala Leu Leu Val Asp Ala

gcc acc ggc cag gtt atc gac gaa ggc cgc gcg agt cac ccg agc ggg 211
Ala Thr Gly Gln Val Ile Asp Glu Gly Arg Ala Ser His Pro Ser Gly

30 35

tcg gag gta gat cca cgt gcg tgg atc gct gcg ctg gat caa gct acc 259 Ser Glu Val Asp Pro Arg Ala Trp Ile Ala Ala Leu Asp Gln Ala Thr

gag ggg ttg tta gaa cgc gcg gac gct gta tct att gca ggc cag cag 307 Glu Gly Leu Leu Glu Arg Ala Asp Ala Val Ser Ile Ala Gly Gln Gln 55 60 65

cac ggc atg gtg gcg ttg gat gaa aac gat gaa atc gtt cgc ccg gcg 355
His Gly Met Val Ala Leu Asp Glu Asn Asp Glu Ile Val Arg Pro Ala
70 80 85

ttg tta tgg aat gac act cgt tct gcc cag gct gcg ttg gat ctc aat
Leu Leu Trp Asn Asp Thr Arg Ser Ala Gln Ala Ala Leu Asp Leu Asn
90 95 100

gag Glu	gag Glu	atc Ile	ggc Gly 105	ggc Gly	gat Asp	cag Gln	gct Ala	gcg Ala 110	gta Val	gat Asp	gcc Ala	acg Thr	gga Gly 115	agt Ser	gtg Val	451
								aaa Lys								499
								gcg Ala								547
								cgc Arg								595
								tac Tyr								643
cgc Arg	acc Thr	gat Asp	cta Leu 185	gct Ala	gcc Ala	ttg Leu	gcg Ala	ctg Leu 190	ggc Gly	cat His	gag Glu	gtg Val	gaa Glu 195	ctt Leu	cct Pro	691
								gcg Ala								739
								aat Asn								787
								gtg Val								835
								gtc Val								883
								gcg Ala 270								931
								ttc Phe								979
								gca Ala				Gln				1027
ggt Gly 310	ggc Gly	gtg Val	acg Thr	ctc Leu	cag Gln 315	cct Pro	tat Tyr	ttg Leu	gag Glu	ggc Gly 320	gag Glu	cgt Arg	acg Thr	ccg Pro	aat Asn 325	1075
cgt Arg	ccc Pro	gca Ala	gca Ala	cgt Arg 330	ggc Gly	gtt Val	ttg Leu	gct Ala	gga Gly 335	cta Leu	aac Asn	tgt Cys	gca Ala	acg Thr 340	acc Thr	1123

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cgc Arg	atc Ile 375	cag Gln	ctc Leu	atc Ile	ggt Gly	ggc Gly 380	ggc Gly	gcg Ala	cgt Arg	tca Ser	cag Gln 385	gcg Ala	gtt Val	cgt Arg	gag Glu	1267
att Ile 390	gcc Ala	cct Pro	gag Glu	att Ile	ttc Phe 395	ggc Gly	cat His	gag Glu	att Ile	gtg Val 400	gtt Val	cca Pro	gaa Glu	ccc Pro	gct Ala 405	1315
gaa Glu	tat Tyr	gtg Val	gcg Ala	ttg Leu 410	ggt Gly	gca Ala	gct Ala	cgt Arg	cag Gln 415	gcg Ala	gca Ala	tgg Trp	gcg Ala	ctg Leu 420	tcg Ser	1363
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Leu	Leu	Val	Asp 20	Ala	Ala	Thr	Gly	Gln 25	Val	Ile	Asp	Glu	Gly 30	Arg	Ala	
Ser	His	Pro 35	Ser	Gly	Ser	Glu	Val 40	Asp	Pro	Arg	Ala	Trp 45	Ile	Ala	Ala	
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Ile 65	Ala	Gly	Gln	Gln	His 70	Gly	Met	Val	Ala	Leu 75	Asp	Glu	Asn	Asp	Glu 80	
Ile	Val	Arg	Pro	Ala 85	Leu	Leu	Trp	Asn	Asp 90	Thr	Arg	Ser	Ala	Gln 95	Ala	
Ala	Leu	Asp	Leu 100	Asn	Glu	Glu	Ile	Gly 105	Gly	Asp	Gln	Ala	Ala 110	Val	Asp	
Ala	Thr	Gly	Ser	Val	Tyr	Val	Ala	Ser	Leu	Thr	Ala	Thr	Lys	Met	Arg	

115 120 125

Trp Met Arg Asp His Glu Pro Glu Asn Ala Ala Arg Thr Ala Ser Val
Met Leu Pro His Asp Phe Leu Thr Trp His Leu Met Gly Arg Gly Arg
145

Val Thr Asp His Gly Asp Ala Ser Gly Thr Gly Tyr Tyr Ser Thr
175

Arg Asp Arg Ala Trp Arg Thr Asp Leu Ala Ala Leu Ala Leu Gly His 180 185 190

Glu Val Glu Leu Pro Glu Leu Leu Ala Pro Asn Ala Ile Ala Gly Thr 195 200 205

Thr Pro Gly Gly Val Lys Val Ala Ala Gly Thr Gly Asp Asn Ala Ala 210 215 220

Ala Ala Leu Gly Leu Asp Leu Gln Pro Gly Asp Val Ser Val Ser Ile 225 230 235 240

Gly Thr Ser Gly Val Ala Gly Met Thr Val Gln His Ser Val His Asp 245 250 255

Pro Ser Gly Leu Val Thr Gly Phe Ala Asp Ala Thr Gly Ala Tyr Phe
260 265 270

Pro Leu Ala Cys Thr Leu Asn Gly Ala Pro Val Leu Glu Phe Gly Arg 275 280 285

Arg Ile Leu Gly Val Glu Trp Glu Glu Phe Asp Ala Leu Ala Leu Ala 290 295 300

Ala Gln Pro Gly Ser Gly Gly Val Thr Leu Gln Pro Tyr Leu Glu Gly 305 310 315 320

Glu Arg Thr Pro Asn Arg Pro Ala Ala Arg Gly Val Leu Ala Gly Leu 325 330 335

Asn Cys Ala Thr Thr Arg Glu Asp Phe Ala Arg Ala Thr Val Glu Gly 340 345 350

Leu Leu Ala Leu Asp Asp Ala Val Thr Ala Leu Val Glu Ala Thr 355 360 365

Gly Val Pro Val Gln Arg Ile Gln Leu Ile Gly Gly Gly Ala Arg Ser 370 375 380

Gln Ala Val Arg Glu Ile Ala Pro Glu Ile Phe Gly His Glu Ile Val 385 390 395 400

Val Pro Glu Pro Ala Glu Tyr Val Ala Leu Gly Ala Ala Arg Gln Ala 405 .410 415

Ala Trp Ala Leu Ser Gly Glu Ala Thr Pro Pro Gln Trp Pro Thr Pro 420 425 430

Gly Ser Asp Pro His Arg Ala Pro Lys Asn Thr Glu Leu Ser Thr Arg 435 440 445 Tyr Ala Lys Leu Arg Ala Ala Thr Gln Gly Trp Tyr

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Ala Asp Ala	Thr Gly	Ala Tyr		Pro Le 185	eu Ala	Cys	Thr	Leu 190	Asn	Gly	
gca ccg gtg t Ala Pro Val 1 195						Gly					624
gag ttc gat of Glu Phe Asp 1 210				_							672
acg ctc cag of Thr Leu Gln I 225	Pro Tyr										720
gca cgt ggc (Ala Arg Gly \				Asn C							768
ttt gcc cga q Phe Ala Arg A	-	-	Gly I	_		-	_	-	_	-	816
gta acg gcg o Val Thr Ala 1 275						Val					864
ctc atc ggt of Leu Ile Gly (290											912
gag att ttc of Glu Ile Phe 0 305	Gly His										960
gcg ttg ggt q Ala Leu Gly A				Ala Ti							1008
acg cca ccg (Thr Pro Pro (Pro G								1056
aaa aac act o Lys Asn Thr (355						Leu					1104
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- Asn Ala Ala Arg Thr Ala Ser Val Met Leu Pro His Asp Phe Leu Thr 50 55 60
- Trp His Leu Met Gly Arg Gly Arg Lys Val Thr Asp His Gly Asp Ala 65 70 75 80
- Ser Gly Thr Gly Tyr Tyr Ser Thr Arg Asp Arg Ala Trp Arg Thr Asp 85 90 95
- Leu Ala Leu Ala Leu Gly His Glu Val Glu Leu Pro Glu Leu Leu 100 105 110
- Ala Pro Asn Ala Ile Ala Gly Thr Thr Pro Gly Gly Val Lys Val Ala 115 120 125
- Ala Gly Thr Gly Asp Asn Ala Ala Ala Ala Leu Gly Leu Asp Leu Gln 130 135 140
- Pro Gly Asp Val Ser Val Ser Ile Gly Thr Ser Gly Val Ala Gly Met 145 150 155 160
- Thr Val Gln His Ser Val His Asp Pro Ser Gly Leu Val Thr Gly Phe 165 170 175
- Ala Asp Ala Thr Gly Ala Tyr Phe Pro Leu Ala Cys Thr Leu Asn Gly
 180 185 190
- Ala Pro Val Leu Glu Phe Gly Arg Arg Ile Leu Gly Val Glu Trp Glu 195 200 205
- Glu Phe Asp Ala Leu Ala Leu Ala Ala Gln Pro Gly Ser Gly Gly Val 210 215 220
- Thr Leu Gln Pro Tyr Leu Glu Gly Glu Arg Thr Pro Asn Arg Pro Ala 225 230 235 240
- Ala Arg Gly Val Leu Ala Gly Leu Asn Cys Ala Thr Thr Arg Glu Asp 245 250 255
- Phe Ala Arg Ala Thr Val Glu Gly Leu Leu Leu Ala Leu Asp Asp Ala 260 265 270
- Val Thr Ala Leu Val Glu Ala Thr Gly Val Pro Val Gln Arg Ile Gln 275 280 285
- Leu Ile Gly Gly Gly Ala Arg Ser Gln Ala Val Arg Glu Ile Ala Pro 290 295 300
- Glu Ile Phe Gly His Glu Ile Val Val Pro Glu Pro Ala Glu Tyr Val 305 310 315 320
- Ala Leu Gly Ala Ala Arg Gln Ala Ala Trp Ala Leu Ser Gly Glu Ala 325 330 335
- Thr Pro Pro Gln Trp Pro Thr Pro Gly Ser Asp Pro His Arg Ala Pro

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Gln Gly Trp Tyr 370

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Asp Leu Thr Ala Lys Val Gln Arg His Pro Glu Pro Gly Glu Thr Leu
25 30 35

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gct gtg gcg gca gcg caa tta ggt gcc aaa gtc acc atg atc ggt gcg 307 Ala Val Ala Ala Ala Gln Leu Gly Ala Lys Val Thr Met Ile Gly Ala 55 60 65

gtc gga acc gat caa atg gct ggc gag gcg ctg aca cat ttg cgt caa 355 Val Gly Thr Asp Gln Met Ala Gly Glu Ala Leu Thr His Leu Arg Gln 70 75 80 85

tca gga gca gat atg tcc gcg att gcc act gtg gac ggt ccc act ggt 403 Ser Gly Ala Asp Met Ser Ala Ile Ala Thr Val Asp Gly Pro Thr Gly

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atc cct ggc gct aac gct tct gtc acc gcg gaa ttt gtt gat aaa cac 499. Ile Pro Gly Ala Asn Ala Ser Val Thr Ala Glu Phe Val Asp Lys His
120 125 130

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